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ADVISORY BOARD ON

RADIATION AND WORKER HEALTH

VOLS. II, III

DAY TWO

ABRWH BOARD MEETING

The verbatim transcript of the

Meeting of the Advisory Board on Radiation and

Worker Health held at the Doubletree Oak Ridge,

Oak Ridge, Tennessee, on January 25, 2006.

<u>C O N T E N T S</u> January 25, 2006

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- MR. MARTIN DELOZIER
- MR. JAMES HACKWORTH
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PROCEEDINGS

(8:45 a.m.)

WELCOME AND OPENING COMMENTS

DR. PAUL ZIEMER, CHAIR

DR. LEWIS WADE, EXECUTIVE SECRETARY

1 DR. ZIEMER: Good morning, everyone. This begins day 2 two of our meeting of the Advisory Board on 3 Radiation and Worker Health here in Oak Ridge. 4 I want to give you the usual reminders, and 5 that is to register your attendance with us 6 today at the registration books out in the 7 corridor. Also, members of the public who wish 8 to address the assembly later today at the 9 public comment period, which is at 7:30 this 10 evening, please register on the book that is 11 also in the -- in the corridor. 12 Again I'll remind you that there are copies of 13 the various documents that are being used today 14 on the table over on my far right. 15 Board members, you should have received this 16 morning copies of the minutes of the August and 17 October meetings, and we will review those minutes tomorrow, but I want to make sure all 18 19 of you have those. 20 The first main item on our agenda this -- well, 21 let me pause here and ask Dr. Wade, our

1 Designated Federal Official, if he has any 2 additional general remarks this morning. 3 DR. WADE: Well, only to -- I mean I -- for 4 some reason I feel compelled to thank the Board 5 for its hard work. There is a tremendous amount of material on your plate, and I know 6 7 that it's difficult and I applaud your efforts. 8 I also applaud your efforts at a level of 9 transparency in work that I've not seen in my time associated with such boards. 10 I think it 11 reflects well on not only the hard work you do, 12 but the way you do that work, and I think a 13 note of thanks is appropriate, so thank you for 14 that. 15 Just to usurp Paul for one minute, on the 16 Pacific Proving Grounds we have no conflicts of 17 interest on the Board, so we are free to do 18 anything we want to do in any way we want to do 19 it. PACIFIC PROVING GROUNDS SEC DR. PAUL ZIEMER, CHAIR 20 DR. ZIEMER: Okay. That opens the door pretty 21 22 wide. Well, I -- that --23 DR. DEHART: Could we visit? 24 DR. ZIEMER: -- that does bring us to the first

main item on our agenda, which is the SEC

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1	petition on the Pacific Proving Grounds. We
2	will have a presentation from NIOSH by Dr.
3	Neton. That will be followed by a presentation
4	from the petitioners, and hopefully we'll have
5	on the line Danella Karo. I don't know if
6	Danella's on the phone yet
7	MS. KARO: Yes, I have just (unintelligible)
8	DR. ZIEMER: Danella, welcome.
9	MS. KARO: thank you.
10	DR. ZIEMER: Please let us know if you have
11	difficulty hearing. So we'll hear then from
12	Danella, and then or is it's Daniella, I
13	guess, is it Daniella?
14	MS. KARO: It's Daniella, yes.
15	DR. ZIEMER: I want to pronounce it correctly.
16	MS. KARO: That's okay. Thank you.
17	DR. ZIEMER: Thank you. And then also present
18	with us this morning is Paul Blake. Paul is
19	here on behalf of the Defense Threat Reduction
20	Agency, and as many of you know, DTRA is
21	involved in a counterpart program involving
22	compensation for atomic veterans. And Board
23	members, I hope you received the recent
24	DR. WADE: It's in the book.
25	DR. ZIEMER: document is it in the book?

DR. WADE: Yeah, it should be everything.
DR. ZIEMER: -- in the tab, as well, with a

number of comments and concerns that were raised by DTRA, and we'll have an opportunity to discuss those, as well, in connection with this petition.

So let us begin with the presentation by NIOSH.

DR. WADE: And Ms. Karo, please let us know if you have any trouble hearing at all.

MS. KARO: Thank you, I can hear you quite well. Thank you.

PRESENTATION BY NIOSH, DR. JIM NETON, NIOSH

DR. WADE: Okay. And now Dr. Neton will start to talk. This is -- this one is proof of the fact that there are no simple SECs.

DR. NETON: Thank you. Good morning, everybody. I'm going to discuss the SE--evaluate-- NIOSH's evaluation of SEC Petition

No. 20, which was -- was received by NIOSH under a different -- a different criteria or different definition. We now are evaluating the Pacific Proving Grounds in total and not just Operation HARDTACK 1, which was the basis for the petition. We'll get into that a little bit later.

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I just want to start -- that's interesting.

The font changed on my slides overnight. It's still readable, fortunately, but that's really odd.

The first slide I just want to show you is a little background information on what we mean when we talk about Pacific Proving Grounds, and the Pacific Proving Grounds is a series of atolls and islands in the Marshall Islands -including Enewetak, Bikini, Johnson Island and Christmas Island -- where a series of nuclear tests were conducted by the military -- U.S. military, starting in 1946 and finishing in 1962. As the slide indicates, it started with Operation CROSSROADS and, through various series, completed with DOMINIC in '62. There were a number of detonations at each of these tests. Each of those detonations had specific names. But suffice it to say that in total there were a little over 100 detonations in the Pacific Proving Grounds during this time period -- of various natures, whether they were underwater, air bursts, surface bursts, that sort of thing. It was a pretty rigorous testing program for nuclear weapons.

As I indicated, the petition as received by NIOSH, initial class, was for all scientists and couriers who were employed at Enewetak Atoll during Operation HARDTACK 1 from July 1st, 1958 through August 31st, 1958 -- a very narrow period of time, a very narrow window. As usual with these petitions, NIOSH takes the opportunity to evaluate beyond the established class to see if there were -- there's any additional exposure scenarios and time periods that should be evaluated and could be possibly covered under this petition.

I'm going to take a similar tack that I have -NIOSH has with past SEC petition evaluations
and sort of go through the process and show you
and highlight the various things that we've
looked at as far as this petition -- and all of
these have been summarized in the evaluation
report that you've received.

So as usual, once a petition meets the criteria it become qualified. This petition was qualified April 11, 2005, and the regulations require that we notify the petitioner and publish a notice in the *Federal Register*. The petitioner was notified and a notice was

published in the Federal Register on May 5th, 2005.

In keeping with our process we evaluate the petition using the guidelines in 83.13, and we submit a summary of the findings of the evaluation report to the Board. That report was sent to the Board on October 20th. It was also sent to the petitioners on October 20th, 2005. A notice that the petition report would be discussed at the Advisory Board, this meeting here, was published in the Federal Register on January 18th, 2006.

There was a supplement submitted to the Board and the petitioners fairly recently, on January 20th, 2006, and that was a result of a couple of things. It was primarily motivated based on a letter that we -- that NIOSH had received from Dr. Paul Blake of the Defense Threat Reduction Agency on December -- it was dated December 22nd, 2005. That letter expressed concern that the petition report contained several misrepresentations of how the -- of a National Research Council report that was published on the DTRA program, the Defense Threat Reduction Agency's programs. And it

also had a discussion of misconceptions that NIOSH may have had about the operating status of the DTRA program. So in this supplement we attempted to address the issues raised in the report, and I'll discuss that in detail a little bit further.

In addition in the supplement we took the opportunity to slightly modify the class definition that we were proposing, and somewhat narrowed the focus of the petition -- of the proposed class, and we'll talk about that later, as well.

Moving on with the evaluation process, as in past petitions there's a two-pronged test established by EEOICPA to determine two things; one, is it feasible to estimate the levels of radiation dose with sufficient accuracy; and secondly, is there a reasonable likelihood that the dose may have endangered the health of the workers. In reference to sufficient accuracy, the regulation states that we need to have access to data that can estimate the maximum radiation dose for every type of cancer that NIOSH would need to reconstruct at this site. And in fact we would need to do this under

plausible exposure circumstances. So in that respect it would not be sufficient -- and in practice we've seen in other SEC petition evaluations -- to, for instance, multiply some doses by a very high multiplier to assure that we bounded the doses. That would not meet our criteria of sufficient accuracy under the regulation.

To determine if we can do these doses with sufficient accuracy, we went about identifying and reviewing a number of data sources of available information to see if we could feasibly do these dose reconstructions. And so in keeping with the hierarchical approach in the dose reconstruction regulation, we went about looking for sources of personal monitoring data, area monitoring data and testing processes and radiation source That is, do we have any bioassay materials. data out there, which would be our choice of preference for doing dose reconstructions. Barring that, do we have any area monitoring samples, air samples, that sort of thing. lacking that, could we possibly reconstruct these doses using a source term analysis.

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1 The resources available were from the usual 2 cast of characters. We have a dose 3 reconstruction database where in fact we have information from claimants. We looked at the 5 claims that we have in our database for claimants from the Pacific Proving Grounds, 6 7 looked -- and searched those for any evidence 8 of internal/external monitoring data, air 9 monitoring data of that sort. We've been 10 successful in the past at finding this type of 11 information on claim files. We've also looked 12 at our research database where NIOSH, with 13 ORAU's assistance, has gone out and conducted 14 extensive data capture efforts to retrieve 15 archived records, looking for information 16 related to these source terms -- bioassay, area 17 monitoring, that sort of thing. 18 We also searched publicly available records 19 that are out there. The Defense Threat 20 Reduction Agency has a very rich web site that 21 contains a number of reports detailing the 22 specifics of many of these tests, as well as 23 some Department of Energy web sites and 24 Lawrence Livermore documents. 25 And lastly, we looked through the documentation and evaluated the documentation or the affidavits that were provided by the petitioner.

We're looking through all these documents and looking at them from the perspective can we do a dose reconstruction and also evaluate these for the -- to determine the basis for health endangerment. That is, were these -- were there sufficient sources of ionizing radiation out there in the environment, and how were these sources of exposure delivered; were they episodic over a period of time, or were they some discrete event where a mere presence would have endangered their health.

As I mentioned, we did review a lot of additional documents that were available on public-accessible web sites, including
Livermore's, DTRA's and DOE's. And in fact we knew that the Defense Threat Reduction Agency was engaged in a similar program to do dose reconstructions for military personnel who were at these test sites. And so we engaged in several conversations with them via telephone and ultimately ended up having a site visit with DTRA contractor personnel who were

1 actually doing these dose reconstructions to 2 evaluate their process to see if it may be of 3 use in reconstructing doses for purposes of 4 EEOICPA. 5 In fact, at the end of our visit we received 6 some -- an example dose reconstruction provided 7 by them for a test case. The dose 8 reconstructions under the Defense Threat 9 Reduction Agency's programs are -- at least the 10 ones that we received -- were -- used the ICRP-11 30 models for internal dose and estimated a 50-12 year committed dose rather than the annual 13 doses that we are -- we use under this program 14 for our IREP calculations. 15 I might add that the dose reconstructions for 16 DTRA -- that the DTRA program itself, the 17 Nuclear Test Personnel Review program, are conducted for non-presumptive cancers. Under 18 19 the military program, all cancers are --20 there's a list of presumptive cancers, much 21 like in our program, and they are actually 22 doing dose reconstructions for the non-23 presumptive cancers. 24 Okay, I'm going to skip this -- that really 25 weird how that changed the font. I'm going to

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skip this slide and go back to it because I think the process will be a little better if I first discuss -- continue on with the evaluation process.

We did look at affidavits and some documentation provided by the petitioners. This consisted of a signed affidavit from the petitioner which discussed that there were a number of scientists gathered in areas where boxes had leaked radiation. There was an indication of civilian personnel who may have been swimming in contaminated waters. there was a contention that there was no monitoring for internal exposure from ingestion or inhalation during these time periods. Supporting that affidavit was a -- some excerpts from a textbook on leukemia that discussed an epidemiologic report done by Caldwell, which showed a statistically significant -- significantly elevated incidence of leukemia at one test shot, I think it was Operation SMOKY. And also it discussed the fact that even -- there were low doses of external exposure, but the epi study was confounded by the fact that there was no

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evidence of any internal monitoring to supplement that dose to do the analysis. The summary of the available monitoring data that we were able to come up with from the sources that we evaluated was that we found that there was considerable external monitoring data. We had evaluated a NIOSH database and we found most people that had filed petitions from the Pacific Proving Grounds -- almost all people -- had external badge results, so there apparently had been a large percentage of workers had been badged. But we found no evidence in our files of any internal monitoring data. And in fact we ran acro-- we encountered almost no internal exposure data for any of the test shots.

There was some indication early on at Operation CROSSROADS that some bioassay samples were taken. However, we could not find these records. And in looking at some of the data that were publicly available, the quality of the data and the reliability of the bioassay data for the CROSSROADS tests were questioned. These measurements were done on board ship with available survey instruments. There did not

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appear to be any long-range plan as to how these things would be measured. There's some evidence of beta activity and things of that nature, but the data themselves do not appear to have a very good pedigree.

There was some off-site air sampling. are primarily environmental samples that were not taken with the intent of reconstructing doses to the civilians. They were more on offsite locations, and although they could be related to the workers -- or the civilians' exposures, we felt that it would be difficult to do so. Or actually we felt there was insufficient data to be able to do that. Okay, let me go back -- let me go back to the DTRA program again. You know, given that we found nothing available for the traditional bioassay sampling that we could reconstruct internal doses, and we felt the area monitoring -- the environmental monitoring data were insufficient, we knew that DTRA was using an approach that related the external dose results on the badge, or some synthesis of the environmental exposures using environmental monitoring external data, to estimate the

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internal dose. In other words, they take the badge reading, and it would be directly converted through this sophisticated computer program into an internal exposure. It's a very sophisticated model, blast -- it uses blastspecific source terms based on the composition of the individual fallout constituents. once the ground-based contamination is assessed, then there's a resuspension factor applied to estimate the internal dose. This had some appeal to us. We did look through it. But in reviewing this whole program we did recognize that the National Research Council did review this process in 2003, a fairly lengthy review was conducted by the National Research Council, and there were a number of findings that were identified that -that questioned the credibility of the upper bounds that were raised. The model themselves -- the model itself was not said to be insufficient or scientifically not valid. just -- it was stated, and I'll quote you one of the things that concerned us in this report. It's stated that the sources of uncertainty -sources and uncertainty of estimating

radionuclide concentrations in depositive fallout based on external photon exposure have not been evaluated, and the reliability of the

methods is unknown.

In other words, they were saying we don't -we're not -- necessarily this is -- this is not
a valid technique, it's just that, you know,
one has not gone and looked at all these
sources of uncertainty and determined if it's
really valid for making these assumptions.
They never questioned that DTRA could not put
an upper bound on this program. In fact, I
would like to state that in our review, we were
evaluating this program only for purposes of
reconstructing doses with sufficient accuracy
under EEOICPA. We were not questioning DTRA's
ability to reconstruct doses for purposes of
their program.

But that finding was one of the findings -there were -- but I should point out, and this
is one of DTRA's concerns, is that there were a
number of findings that indicated that doses
could be either underestimated or
overestimated. Now our evaluation report
focused on the overes-- or underestimating

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conditions, but -- because we have to be able to put an upper bound -- there is no real -there was really no real indication in the report, though, about the magnitude of the corrections in either direction -- or the uncertainties in either direction. So it's not obvious to us from reading the report that these things -- these countervailing factors would cancel each other out and indeed the dose reconstructions are sufficiently accurate. DTRA did put together a plan of action and in June of 2004 it indicated that they were going to evaluate these issues and complete their analysis in a two-year time frame, which would put their completion date somewhere around June 2006 -- possibly six months or so from now. Interim guidance was issued in July 16th by DTRA that allowed them to move forward with dose reconstructions. And in that interim quidance, which is attached to the letter that Dr. Blake sent NIOSH, it indicates that they believe that if one multiplies internal doses by a factor of ten, they would be sufficiently bounding and put a credible upper bound on the doses and would allow them to move their dose

reconstructions forward. We've not seen any scientific analysis of that, and until such time as we get a handle -- at least NIOSH feels -- on the magnitude of the individual sources of uncertainty, the fact of multiplying the doses by a factor of ten would not constitute sufficient accuracy under the requirements of our program.

So in looking through all of the data -- the available data and the DTRA program and the evaluation report supplement, we revised the class definition at Pacific Proving Grounds from covering Operation HARDTACK 1 to the definition you see on the screen, which is all employees of DOE, DOE contractors or subcontractors employed at the Pacific Proving Grounds from 1946 through 1962 who were monitored or should have been monitored for exposure to ionizing radiation as a result of nuclear weapons testing at the Pacific Proving Grounds.

We did find evidence that sources of internal exposure existed from these multiple detonations of weapons, but the exposures to the civilians were a result of contamination

caused by inhalation of fallout from the nuclear detonations, not from the direct exposure to the criticality event that occurred when the weapons were detonated. And we believe that, in looking at all the records, we currently lack access to sufficient bioassay or air monitoring data to estimate doses associated with inhalation to radionuclides at the Pacific Proving Grounds.

We further believe that health was endangered. We cannot estimate these doses with sufficient accuracy, and the evidence indicates that

We cannot estimate these doses with sufficient accuracy, and the evidence indicates that workers may have accumulated internal exposures through episodic intakes of radionuclides; that is, from the resuspension of fallout and the direct inhalation of fallout from the weapons tests. These are internal doses. By nature they are defined as chronic exposures and not acute exposures, so therefore we -- we are not suggesting that mere presence at this site would constitute membership in the class. But 250 days, as defined in the regulation, would constitute membership.

And this is the proposed class definition again, and a summary. We're summarizing the

1 class as 1946 through 1962, it's not feasible 2 to reconstruct doses with sufficient accuracy, 3 and we believe health was endangered. 4 With that, I would be happy to answer any 5 questions. Thank you very much, Jim. Let me 6 DR. ZIEMER: 7 begin with a question on a slide, and I'm not 8 sure you actually put this one up. It's in our 9 book. The NIOSH claims tracking system 10 indicates there are 65 cases that would be in 11 this class. Is that number still correct -- in 12 the class as redefined? 13 DR. NETON: Yeah, that's right. I must have 14 skipped right through that one in my... 15 DR. ZIEMER: Now I'm curious about the next 16 line where it suggests that you actually have 17 completed some dose reconstructions, which 18 seems somewhat counter to what you said, so --19 DR. NETON: That's correct. 20 DR. ZIEMER: -- just explain to us what that's 21 about. 22 This is not -- this has DR. NETON: Yeah. 23 happened with other SECs where, you know, we 24 have moved cases forward where we thought 25 possible. In reviewing the SEC petition

evaluation -- SEC petition, though, we've come to the conclusion that the ones that we've processed and moved forward have not been done with -- reconstructed with sufficient accuracy. This has happened at Y-12. In the past when we've added classes we've actually moved a few cases out, but in looking at the bigger picture and the definition of sufficient accuracy, we've come to that conclusion.

DR. ZIEMER: Thank you. My next question has to do with the information sources. And I noticed in the DTRA document -- I suspect maybe Dr. Blake will speak to this, but there's a suggestion that there are additional data sources that NIOSH failed to look at. Could you speak to that issue?

DR. NETON: Yeah, there was a suggestion that there were some data sources available -- I think in particular it was related to the bioassay data themselves, related to some medical files and some Naval transmissions and that sort of thing. We have not seen those data sources. They speak to the internal bioassay data that we were looking for. And in fact, in talking with the DTRA program

1 personnel, at least the contractor personnel, 2 they don't use those pieces of information 3 either in their program. Those are -- they 4 solely reconstruct doses based on the external 5 badge results. But we have not actually seen 6 the bioassay records. 7 I did mention that the CROSSROADS samples were 8 taken early on, done on board a ship, and I 9 think that the program itself had some 10 questionable pedigree as to how -- how well 11 they could measure what they were trying to do. 12 DR. ZIEMER: Is there any indication that those 13 sources would be readily available in a 14 reasonable time period that would allow one to 15 at least examine them for usefulness, or... 16 DR. NETON: I'd actually defer to Dr. Blake on 17 that question 'cause the --18 DR. ZIEMER: Maybe when Dr. Blake speaks to --19 This is fairly new information that DR. NETON: 20 we received -- yeah. 21 DR. ZIEMER: Yeah, I understand. We just -- we 22 just got the document I think a couple of days 23 ago. 24 Okay, Dr. Melius and then Mark. 25 DR. MELIUS: Yeah, Jim, I have a question.

don't know whether it may be a question that

Department of Labor will need to answer, but I

believe this is the first time we've dealt with

a -- hear a SEC definition involving a

"monitored or should have been monitored", and

I'm just -- like to understand how that

determination is -- is made. I believe, if I

understand the process, once -- if this were

accepted, et cetera, that the Department of

Labor would be making that determination?

DR. NETON: That's correct.

DR. MELIUS: But I'm trying to understand here how that -- how that (unintelligible). Is that done by job classification in some way or job description, or what is...

DR. NETON: Yeah, I might have to defer the answer to that question to Department of Labor, but the decision to narrow the class to workers who were monitored or should have been monitored was based after some discussion with the Department of Labor and to focus the membership in the class on those who were actually exposed. If one does not have that proviso there, then people who were never even working in the presence of radioactive

1	materials that were employed working on the
2	Pacific Proving Grounds project, for example,
3	would be eligible.
4	DR. MELIUS: Yeah, I and that part I
5	understand. I'm just trying to understand how
6	you make it operational how you make that
7	determination 'cause particularly in this
8	situation where it might not be straight at
9	least not it doesn't seem straightforward to
10	me, that's all.
11	DR. ZIEMER: Here's Pete
12	MR. TURCIC: That is the
13	DR. ZIEMER: Turcic from Labor.
14	MR. TURCIC: That is the provision. That exact
15	provision is in all of the Congressionally-
16	mandated SECs.
17	DR. MELIUS: Correct, and that's why
18	MR. TURCIC: Yeah, and the way we handle that
19	is by occupation.
20	DR. MELIUS: Yeah.
21	MR. TURCIC: At Amchitka the way we handled it
22	was by policy. We were able to identify that
23	there was ionizing radiation after the first
24	shot, so then presence after the first shot.
25	At the other sites it's the was badged or

1 should have been badged applies to what does 2 the policy -- what would a policy now --3 DR. MELIUS: Yeah, okay. 4 MR. TURCIC: -- and then work backwards from 5 there. DR. MELIUS: Okay. No, it just -- I realized 6 7 it was in the Congressionally-mandated SECs, 8 but we -- we -- it's never come up here and 9 we've never discussed it and I'm just trying to 10 understand how it would work. Thank you. 11 MR. GRIFFON: Jim, I was just wondering, you 12 mentioned -- either in the presentation or in 13 your written statements -- that you have 14 reviewed some cases, some DTRA cases? DR. NETON: Just -- well, there was one case we 15 16 were provided a dose reconstruction. 17 MR. GRIFFON: Oh, just one case. Okay. And 18 did it include internal dose --19 DR. NETON: Yes. 20 MR. GRIFFON: -- estimates? 21 DR. NETON: Yes. 22 MR. GRIFFON: And those were done -- I mean... 23 DR. NETON: They were internal dose estimates 24 based on -- I don't recall if it was from the 25 badge reading or the estimated, you know,

1 external exposure from environmental conditions 2 or survey measurements. But nonetheless it 3 would take the external result and estimate the internal result from -- from the external and 4 5 report -- I think it's -- I'm -- it's a 50-year 6 committed dose to the organ as a result of that 7 exposure, so... 8 MR. GRIFFON: And it's your feeling that the 9 models that they used or are currently 10 implementing couldn't credibly bound doses for 11 the claimants in this population? Is that... 12 That's correct, it could not bound DR. NETON: 13 the internal exposures based on the -- based on 14 the current -- the NRC review that was done 15 that questioned the reliability and the 16 uncertainty of the method that had not been 17 demonstrated, we are not certain that the 18 exposures that are calculated for internal dose 19 at this point for this program are sufficiently 20 accurate. 21 DR. ZIEMER: Roy DeHart. 22 DR. DEHART: Jim, I'm curious about the numbers 23 that we're seeing up there, 65 case 24 definitions. Yet over the period of time I 25 would assume that there would have been perhaps

1 1,000 civilians all told through there. 2 there been any active program at this point in 3 time to notify or contact prior workers? 4 DR. NETON: Not to my knowledge. 5 DR. ZIEMER: Any additional questions for Jim? 6 (No responses) If not, thank you, Jim. 7 8 PRESENTATION BY PETITIONERS, DANIELLA KARO 9 Then we will hear from the petitioner, Daniella 10 Karo. And incidentally, Daniella, I'm not sure 11 what the time is out there but I know it's very 12 early in the morning. 13 MS. KARO: Well, actually it's just, you know, 14 past 6:00 a.m. here, 6:18, actually. 15 DR. ZIEMER: Okay. Well, we appreciate your --16 MS. KARO: Well, I appre--17 DR. ZIEMER: -- early morning (unintelligible) 18 in being with us. 19 MS. KARO: Well, thank you for allowing me --20 am I -- do I have the floor? 21 DR. ZIEMER: You may make your presentation 22 now, if you would, please. 23 MS. KARO: Well, I don't really have a 24 presentation, per se. I understand there is --25 I read through the -- through the findings, you know, to my understanding. I realize that

NIOSH has evaluated the data sources and they
were not able to come up with any sources that
were reliable enough to -- to make a
determination about the inhalation and
ingestion, et cetera. And then there's this
very elegant discussion about the various
formulas that DTRA is using and so forth. I
cannot really ask too many questions about this
because that is definitely not my area of
expertise.

But I do have a question to ask because perhaps

maybe I need a little bit clarification on this. I noticed that NIOSH, in their summary of their findings, seems to be bound by the requirements that the workers were employed in the Pacific Proving Grounds or in any other places have to have established a number of work days aggregating at least 250 days worked there within the parameters established for the class. And I know -- apparently this is -- Congress -- that decided to -- to limit, you know, to 250 days, and apparently it specified originally that there were a class of employees that were included in the SEC and this were the

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employees at the -- that were employed at the gaseous diffusion plants and that's where they (unintelligible) at least 250 days before February, 1992. On the other hand, they have allowed employees who were employed on the -on Amchitka Island and were exposed to ionizing radiation, and yet they were not required to have an aggregate of 250 days. So can you address some -- you know, some of these issues? I'm concerned about this upper limit of 250 days because how did Congress come up to this number, an aggregate number? What happens to individuals who were there for several weeks or let's say an aggregate of 150 days. How are those people treated and why is an upper number required in order to establish this kind of a class, and yet there is an exception on the Amchitka? These are my questions.

DR. ZIEMER: Very good question, and we will ask NIOSH to address that. Let me mention that one might argue that in -- at the test site that the episodic nature of some of these might argue for a different or modified approach to that, as opposed to a workplace where there might be chronic exposures. But there is some

rationale nonetheless for the 250, which itself
has a kind of built-in arbitrariness, but is
related to other sites where that's been the
criteria based on Congressional mandates. But
I'll let Jim speak to the issue particularly

here.

DR. NETON: I thought you did very well, Dr.
Ziemer.

I can't -- this is Jim Neton. I can't speak to the legislatively-created Special Exposure Cohort requirements for Amchitka Island. really just -- I have no idea as to why presence was included at Amchitka and why 250 days was required at the gaseous diffusion plant -- I mean why it was different than the gaseous diffusion plants. I can say that within our regulation the 250-day requirement is there for essentially what you would consider almost chronic-based exposure, not something that happened as a discrete event that endangered the health of the employees within a short time frame where they would experience very large exposures. And I think the example provided in the regulation is such as at a criticality event or incident.

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Now one would argue that a nuclear detonation is a criticality incident, but these people were removed from the blast itself. I think there was some exclusion zones around the blast. And their exposures by nature were chronic exposures to fallout that had either deposited -- either inhalation of the fallout or from resuspension of the fallout that had been deposited on the ground, incorporated in their bodies, and the dose is delivered over a -- a time frame, a chronic time frame, not over a very discrete period. So that's the basis for our ruling -- or our decision, rather, that the 250-day criteria applied. I see. I understand what you're --MS. KARO:

when you're talking about the chronic exposures versus the episodic nature, and yet -- I mean somewhere in the report we're talking -- and I think there was a reference to -- particularly to a shot I believe in Enewetak, and may have been the very last one, I'm not so sure, where -- I'm trying to find my notes here -- in fact the (unintelligible) at Enewetak Island apparently was closed for one day because of the fallout from an atmospheric test at Bikini

Atoll designated for -- that was in May 12, and you're talking about resuspension -- so let me ask you a question. So you're saying that basically an individual had to be chronically exposed in order to be affected while the episodic nature kind of removed -- I mean basically the individuals who were episodically exposed may not have had any, you know, exposures that may provoke any cancers. That's the way I understand it.

DR. NETON: I'm not sure I'm following your -MS. KARO: Well, my feeling is that you need a
chronic exposure -- we're trying to define how
we -- you know, Congress came up with this, you
know, 250 days. And so individuals -- it
sounds like you need individuals to be
chronically exposed in order to ultimately, you
know, come down with a cancer versus just an
episodic nature, which there is a likelihood
that the individual would not have that kind of
effect.

DR. NETON: I think I agree with what you're saying in the sense that one -- the dose that's delivered to devel-- to produce the cancer, if it is received over a period of time from a

episodic nature -- and I don't necessarily say it's chronic, but just over a period of time -- one needs to accrue that dose to endanger the health, then the 250-day criteria would apply. If the dose was received in a very short duration, in a discrete incident, then the -- you know, the presence or the time frame would be reduced to whatever that discrete time frame was. But by nature, again, these are internal exposures that are delivered over a period of time.

MS. KARO: And is Amchitka Island an exception?

DR. NETON: Amchitka Island was legislatively added by Congress, and yeah, it's -- it's different than what we're -- it's different than the regulations that we're working under for this program.

MS. KARO: And the reason that they were excluded from this 250 days was because?

DR. NETON: I can't answer that question.

MS. KARO: I see. That was my concern, because I'm sure that there would be individuals who probably were not there, you know -- or some individuals may not have been there for an

aggregate of 250 days, and my concern is how

1 are those individuals going to be treated, on 2 an individual basis or how is -- how are their 3 petitions going to be treated, if indeed this 4 is not -- you know, if this regulation -- this 5 rule is going to be maintained and -- or 6 applied? 7 DR. NETON: I think, as with other SEC classes, 8 if a claimant does not -- or a case does not 9 have 250 days aggregate exposure, then they 10 would be referred to NIOSH for dose 11 reconstruction. 12 MS. KARO: And (unintelligible) --13 DR. ZIEMER: What he's saying is there could 14 still be an attempt, in an individual case, to 15 do a dose reconstruction. 16 DR. NETON: That's right. 17 DR. ZIEMER: May -- may or may not be feasible, 18 though. 19 DR. NETON: We would do the dose 20 reconstruction, given the techniques that we 21 have available that we believe are sufficiently 22 accurate. In this case we're suggesting that 23 we can't do internal -- we -- at this present 24 time we do not have a technique to do dose --25 internal dose reconstructions with sufficient

accuracy.

MS. KARO: And how is this going -- if an individual is going to be considered outside of a class, I suppose if this goes -- you go ahead and accept this as a class, if an individual does not meet that upper requirement of 250 days -- aggregate days, then my understanding is you would be doing it on an individual basis. And yet -- and yet how will you be able to establish, you know, the dose exposure if you don't have, you know, fully access to -- to internal exposure, as well?

DR. NETON: That's a very good question. We would attempt to reconstruct the doses that we knew we could. And in this case, we believe that the external dosimeter results are -- are acceptable or sufficiently accurate for our purposes of reconstructing doses, and we would attempt to do an external dose reconstruction. But we would not at this time, if this class were added under these conditions, be able to do internal dose reconstructions.

MS. KARO: And that's only for the people who were in the class, but what about when you have to -- to deal with them on a one by one basis

1 because they did not meet that upper 250 days? 2 Since you don't have a formula yet that would 3 allow you to figure out the internal dose, then 4 how are you going to, you know, treat those on 5 an individual basis then? 6 DR. ZIEMER: Perhaps the --MS. KARO: How will you be able --7 8 DR. ZIEMER: -- question is if they don't --9 MS. KARO: -- to actually establish with some 10 accuracy the -- the dose exposure? 11 DR. ZIEMER: Yeah, perhaps it's -- the question 12 of if you have a case where you say actually we 13 can't reconstruct this dose and the 250-day 14 criteria is not met, what happens to that case, 15 is --16 MS. KARO: That's correct. 17 DR. NETON: Well, all we're saying at this 18 point is we cannot accurately reconstruct the 19 internal dose. We would reconstruct the 20 external dose to the extent we could, and we 21 would use -- use that as a basis for our dose 22 reconstruction. But we would not be able to 23 add any internal dose because we don't have 24 tools --

DR. ZIEMER: And if the --

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1 DR. NETON: -- to do that at this time. 2 DR. ZIEMER: If the person didn't meet the POC 3 criteria based on external, then the case would 4 be denied then --5 DR. NETON: That's correct. 6 **DR. ZIEMER:** -- apparently. Additional 7 questions, Daniella? 8 MS. KARO: No, that's -- that's all I have. 9 BOARD DISCUSSION/RECOMMENDATION DR. PAUL ZIEMER 10 DR. ZIEMER: Then let us hear from Dr. -- or 11 let me see if --12 MR. GRIFFON: I was just going to ask Jim --13 Jim, can --14 DR. ZIEMER: Yes, Mark. 15 MR. GRIFFON: -- you describe the nature of --16 of the work for these employees? I mean -- you 17 know, it seems like we're looking at this 18 (unintelligible), you know, they were there 19 during the fallout, I guess, but they probably 20 went back in and did recovery operations or --21 DR. NETON: Yeah, I've looked at some of the 22 cases that we have in our files, and it's all 23 over the map. There are these scientific 24 couriers who were there delivering documents

and materials and things of that nature, all

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1 the way to people who were operating land-2 roving equipment, driving all over the -- the 3 islands and that sort of thing for -- for a 4 long -- long periods of time, reconstruction 5 activities, sort of construction activities, so there -- it covers the waterfront of different 6 7 types of activities, from what I can see so 8 far. 9 DR. ZIEMER: Dr. Melius, another question? 10 DR. MELIUS: No, that was actually my question. 11 DR. WADE: Jim, I would --12 DR. ZIEMER: Lew Wade. 13 DR. WADE: -- just like to maybe add some --14 this is Lew Wade -- some clarification to the 15 petitioner's question and -- and the answer. 16 And please correct me if I'm wrong. What I 17 hear you saying is that it is your belief that 18 the people who might be considered for this 19 class were, in our opinion, shielded from the 20 high radiation associated with the particular 21 blasts. We -- we believe that they would have 22 been removed or in some way shielded from that 23 radiation. 24 DR. NETON: I wouldn't necessarily use the word 25 "shielded", but removed --

1 DR. WADE: Removed. 2 DR. NETON: -- is probably a better -- a better 3 term. 4 DR. WADE: And then they would be exposed then, 5 after the blast, in their work that they would do and that's why we consider the -- the 250 6 7 days as appropriate. 8 DR. NETON: Correct. 9 DR. ZIEMER: In fact, I might add that in the 10 other episodic cases such as the Oak Ridge 11 criticality accident, the main issue was the 12 external dose, so you wouldn't specify someone had to be there 250 days. If they were in that 13 14 room when the excursion occurred, then they 15 were qualified. 16 DR. NETON: That's correct. 17 PRESENTATION BY DTRA, DR. PAUL BLAKE 18 DR. ZIEMER: Now we'll hear from Dr. Paul Blake 19 with DTRA. Dr. Blake, welcome. 20 DR. BLAKE: Dr. Ziemer, members of the Board 21 and interested parties, thank you for affording me this opportunity to address in the next five 22 23 minutes -- briefly, I hope -- how the 24 Department of Defense currently generates dose

reconstructions with regard to the Pacific

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Proving Ground, and our plans for the future.

I serve as the program manager for the Nuclear
Test Personnel Review program at the Defense
Threat Reduction Agency. The Defense Threat
Reduction Agency, otherwise known as DTRA, is a
combat support agency of the Department of
Defense. DTRA functions as the executive agent
for DOD supporting radiogenic disease claims
brought forth by former DOD personnel who
participated in atmospheric nuclear weapons
testing from 1945 through 1962, served as U.S.
occupation forces of Hiroshima and Nagasaki, or
prisoners of war in the vicinity of Hiroshima
and Nagasaki when the detonations occurred in
1945.

DTRA supports the Department of Veterans
Affairs and, to a lesser extent, the Department
of Justice in evaluation of radiogenic disease
claims. The NTPR program has been in existence
since 1978, and has accumulated a wealth of
documentation associated with these nuclear
events, and developed methodologies for
generating dose reconstructions associated with
these events.

Generating dose reconstructions for atomic

veterans is technically challenging. This fact was extremely well-documented in a recent National Academy of Sciences National Research Council report published in 2003 that Dr. Neton referred to. It's entitled "A Review of the Dose Reconstruction Program of the Defense Threat Reduction Agency".

My agency has been busy in responding to the challenges noted in this publication. We have modified our procedures to overcome these challenges. In fact, at the next meeting of the Veterans Advisory Board on Dose Reconstruction, a board very similar to yours in function but mandated by Congress to oversee the programs that I serve as program manager, I'll be reporting on the status of that progress.

One of the consequences of the NRC report was for the Department of Veterans Affairs to return to DTRA over 1,200 previously-generated dose reconstructions and request that we rework these dose reconstructions, correcting the challenges noted in the 2003 NRC publication. DTRA is in the process of responding to this request.

However, my agency is greatly concerned that the dose reconstructions be performed in a timely manner since many of our veterans are quite elderly and do not have many years to live. Consequently we have made a conscious decision to immediately address these concerns, modify our procedures and begin releasing revised dose reconstructions while the formal process of publishing articles concerning our revised techniques in peer-review journals and releasing revised and new DTRA technical reports on our public web site has preceded us at a slightly slower pace.

This decision has led, in my opinion, to the

This decision has led, in my opinion, to the current situation being discussed today.

Namely, one government agency, NIOSH, has determined that it lacks sufficient information to estimate internal doses associated with Pacific Proving Ground personnel, which a different federal agency, DTRA, is currently in the process of performing this function.

It is my expectation that this problem should be resolved by the end of calendar year 2006.

DTRA, via its contractors and contract staff, has drafted numerous technical reports in

response to the problems noted in NRC 2003. In the next few months papers and reports will begin to be published, and we expect the majority of these will be released by the end of this calendar year.

On another note, I would point out that many of the internal dose challenges noted for the Pacific Proving Ground become even more challenging when addressed for the Nevada Test Site, specifically the scenario of the nuclear detonation blast wave resuspending radioactive fallout from previous detonations, complicating the determination of inhalation dose.

In conclusion, both NIOSH and this Board are in a difficult position. You need to make a timely decision, but you are faced with the current dilemma of two federal agencies proposing two very different solutions for personnel who potentially inhaled radioactive material while performing their duties during atmospheric nuclear weapons testing at the Pacific Proving Grounds. It is my hope that this challenge will be resolved by the end of this calendar year.

And with that, I'd like to open up -- if you

have any questions.

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BOARD DISCUSSION/RECOMMENDATION DR. PAUL ZIEMER

DR. ZIEMER: Thank you, Paul. Could you -- aside from the issue of whether one can reconstruct those doses or not, does DTRA have a 250-day requirement on the individuals for whom you reconstruct the dose?

DR. BLAKE: Well, the requirements actually come from both the Department of Veterans Affairs and Department of Justice with regard to verification of -- to -- are they participants at that place. And these were mandated by Public Law. They're published in the Code of Federal Regulations. But they are different than your 250-day program. They're much more of an episodic nature. The way they're laid out is per -- it differs only slightly between the Department of Justice and Department of Veterans Affairs Code of Federal Regulations. They are defined for each atomic -- atmospheric atomic test, and there's a certain period of time that, if a person was in that area, then they qualify.

What we do in the Department of Defense is we research the records -- and the military kept

very good records back then -- and we pull up and verify, at least in the case of the Department of Veterans Affairs, if that person is a participant. And dependent upon the type of radiogenic disease they have, they either qualify for presumptive compensation or, in the -- we have basically just primarily two cancers now that we do dose reconstructions for, the non-presumptive nature. It's primarily prostate and skin cancer. And so almost all the rest of the cancers are then presumptively compensated through the Department of Veterans Affairs.

DR. ZIEMER: Then is presence at, for example, as little as one test -- would not exclude one from being considered. You would basically look at whatever dose you calculated for that person?

DR. BLAKE: And that's exactly correct. What we find in the case of the Pacific Proving Ground is most of the military personnel involved were part of the Navy, especially those early tests like Operation CROSSROADS. The reason we did those tests was to look at weapons effects. And so the military put a lot

Of people in place where Atomic Energy
Commission, the group that you're looking at,
had a lesser number of people. Our total
population is on the order of about -- when we
look at both Nevada Test Site and Pacific
Proving Ground, about 250,000 people, some of
which are now deceased. But many of those
sailors, for instance at the Pacific Proving
Ground, were on a ship and were there for only
a month or two before they departed. So they
certainly would not have qualified for the 250day figure that's being discussed here.

DR. ZIEMER: Can you give us some idea of the extent to which the individuals who qualify were -- are the doses -- do they tend to be primarily external doses? Are these -- I'm really getting at are these -- are the veterans -- do they tend to be in closer than our workers, or maybe we don't know that, but would they tend to have a bigger contribution from external dose, or do you know what the relative contributions are, by your calculational methods, external versus internal?

DR. BLAKE: By our calculational methods, in general the external dose is the larger of

(unintelligible) --

DR. ZIEMER: The driver.

DR. BLAKE: The driver. There are a few exceptions where internal dose is -- is a concern. But the challenge that Dr. Neton noted and that we have to over-- we have to dwell with is that internal dose is -- is tough to reconstruct based on the data that we have from 50 years ago, and so the uncertainties are normally larger with that component than the external dose.

DR. ZIEMER: And what I'm trying to get at is if the external is the driver, given its uncertainties, it may be, in the DTRA case, that the internal, even -- even if it's not well known, has very little impact. Whereas in our case, if our workers are driven by the internal dose, that may become an overriding issue in knowing that value with a degree of accuracy that become much more important. I'm -- I'm not sure that's the case. I'm kind of asking, is there a difference in how the veterans were positioned and stationed during these tests versus these other workers? Did they tend to be forward, for example, and

1 therefore subject to the external exposures 2 more than these workers? Do we even know that? 3 I think -- I would think we would, but... 4 DR. BLAKE: I can address that just roughly in 5 that many of our veterans were Navy personnel on board ship close to the tests. I think -- I 6 7 believe more of the Atomic Energy Commission 8 personnel were back on the actual islands. 9 There were military personnel back there -- for 10 instance, communications people and so forth --11 too, but probably the majority of veterans were actually on ship for -- for this testing. 12 Uh-huh. So there may be some 13 DR. ZIEMER: 14 differences in -- in how these populations were 15 exposed that might argue for perhaps some of 16 the reasons that we may have to approach them 17 differently, is all I'm saying. I'm concerned 18 myself about two agencies sort of having 19 different answers or different solutions, but 20 maybe there's a reason for that. 21 Dr. DeHart. 22 DR. MELIUS: Paul, can I ask, if it's okay -- I 23 have a sort of follow-up question to that that 24 I'd like to get clarified. And that is, are we 25 really dealing with sort of a mixed group, at

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least on our side, some of whom will have the char-- characteristics of exposure of the -the Navy, the military personnel, and some of which will have other -- you know, another -you know, sort of a longer term exposure that would be different and -- and -- and do we really need to think about how do you separate those two groups out or at least make sure we address both those groups appropriately. some group may be more appropriately addressed through sort of the -- the veterans program and the other sort of addressed looking at sort of this chronic exposure maybe where internal exposures would be more -- more important. that's what I'm trying to wrestle with is -- is understanding sort of what's the right way of navigating this and -- and at the same time that balance that you just said, so --DR. ZIEMER: It's the other side of the same question, and I think that unless one has that kind of specificity in the information database, it would be very hard to separate that out. Any comment on that, Paul? DR. BLAKE: I can't comment too effectively 'cause I'm not really familiar with the Atomic

Energy Commission personnel, sorry.

DR. ZIEMER: Okay. Roy DeHart.

DR. DEHART: My question is parallel to that, basically. I am reasonably certain you had Naval personnel on the ground doing contractual activities -- Seabees and others who were doing things along the atoll or islands -- similar to the exact work that the civilians were doing. They got out of the way when you got ready for the blast. Then they went back in and did whatever they did, creating dust and various other things. So how are you handling those civilians -- or those military who were on the island -- not on the ships -- and would have been extracted to some degree before an explosion?

DR. BLAKE: We do the calculations on -- that were addressed here with regards to internal dosimetry, but primarily it's due to radioactive fallout, usually in almost -- most of the cases. We do look in every -individual case whether there is concerns about instantaneous radiation from the actual nuclear detonation, but it -- in the vast majority of cases our military personnel at the Pacific

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1 Proving Ground were far enough away -- in the 2 case a mile or two -- that there was no 3 radiation from that instantaneous explosion. 4 And so radioactive fallout becomes the greatest 5 concern. And there were a few instances in the Pacific Proving Ground where the winds shifted, 6 7 went the wrong way, and we actually had some 8 acute skin burns where -- and there's a classic 9 case, of course, of a Japanese fishing ship 10 that occurred. But some U.S. service personnel 11 at some radio communications stations also 12 received some significant doses. Other personnel who've received some 13 14 significant doses, a little different than your 15 question, sir, but were some of our pilots and 16 people in aircraft that actually through --17 flew through the radioactive plumes to pick up 18 data. But we certainly did have people on the 19 islands doing some similar, I believe, 20 exercises in locations that probably the Atomic 21 Energy Commission personnel were at, too. 22 DR. ZIEMER: Now Mark. 23 MR. GRIFFON: I just have a que-- I think you 24 sort of answered this question. There is some 25 statements in -- I forget if it's your

statements or Dave Kocher's statements on the
- the internal dose and basically indicating

that some of -- of these -- some of the

findings and the -- and how to resolve those

findings have not been implemented at this

point -- or fully resolved, I guess. And in

the interim -- I think, if I'm reading this

right. In the interim, you have some of these

rules of thumb that you've put in place. Is

that correct?

DR. BLAKE: We call --

MR. GRIFFON: And then -- and then by the end of 2006 you're expecting to have answer or -- or full resolution to -- to these issues.

DR. BLAKE: The NRC report really pushed for doing full uncertainty analysis on our doses, and obviously that -- it would be the preferable method to do. The other approach, though, is to calculate a maximum radiation credible dose, and when we do our doses we calculate out to a 95 percent credibility limit on our calculations that we then forward over to the Department of Veterans Affairs to do the probability of causation calculation with similar software to what this program uses.

What we ended up doing because of these large uncertainties was basically assigning a factor that we believe would be at least at the 95 percent or greater. A constant we try to apply, give the benefit of the doubt to the veteran. So in some cases where we may not be able to say exactly the uncertainty is at the 95 percent confidence level, we actually do -- we -- we argue that we put in place a value that may in fact exceed that.

Where the challenge has occurred with NIOSH is that we've moved ahead -- implemented these procedures, but we've not published them publicly on our web site. And if you're a different federal agency looking at the procedures, you look at some of our publications -- and one was alluded to on how we calculate internal dose based on the radioactive fallout. That publication is from 1985. We're in the process of releasing a revised one here in the next few months. We've gone through four to five major software revisions on that particular code, bringing in place ran-- values up from like ICRP-30 to ICRP-66 and 67 on inhalation dose and so forth,

modernizing both the hardware and software.

But it is a sophisticated code and -- and a lot of assumptions have to go into it because of the circumstances we are under.

MR. GRIFFON: And I guess the -- the rules of thumb, particularly this factor of ten that's sort of mentioned in those statements, this -- this -- there is statistic basis to that --

DR. BLAKE: Yes --

MR. GRIFFON: -- (unintelligible) gets you up near your 95th (unintelligible).

DR. BLAKE: We also have some other publications that are -- that'll be published shortly where we've done bounding analysis on those values. Do we want to move away from those factors, that's the reason we called them interim guidance? Yes, we do. But we had to do that, concerned with our veterans that we had to move ahead with our dose reconstructions, too. And we believe we gave conservative factors at that time. But you'll be seeing shortly other publications that our contractors have worked on where we looked at these concepts of fractionation and -- and bounded them and believe, in most cases, that

1 this is a good figure. 2 DR. WADE: I have a comment to make here. 3 Well, first I'd like to thank you for being 4 here, and I'd like to thank you and your agency 5 for being so professional through this behavior. You've been forthcoming and you've 6 approached this issue in a way that I think 7 8 brings honor on all of us that are federal 9 employees. I think you do high quality work 10 and I think no one disputes that. 11 As a federal employee in this case I'm not 12 embarrassed by the fact that two federal 13 agencies take a different position. What -- I 14 mean that's the way it happens sometimes when 15 you start at different points and move towards 16 a common objective. 17 I am heartened by the fact that this debate is happening in a very open forum and we're 18 19 allowing the issues to be vetted fully. 20 thank you for your participation in that and 21 congratulations to your agency on its program. 22 DR. BLAKE: Thank you, sir. 23 DR. ZIEMER: Okay, Board members, any 24 additional questions? We want to give a full

airing to any of the issues, issues that are

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raised, I think particularly by DTRA in terms of this difference between the agencies and some differences in how things are calculated. But there are some even legal differences in our laws that mandate certain things, so it's -- it's not a bad thing that we're not exactly parallel. We are trying to work together and work closely on approaches as we move forward, but this is one where, in part, there's also a timeliness issue. It may be that one could say well, let's wait six months till they come out with their new guidance or whatever time it is, and you have that issue is somewhat like we had at Mallinckrodt, that there's -- it's a moving target. So one of the timeliness factors I think does come into play and we have to think about that in terms of action, or lack thereof. Henry, and then Richard.

DR. ANDERSON: I guess the question I would have is it sounds like there are some methods that have been developed, they just haven't been made public or they're not completely polished, and I guess my question would be is it possible for those contractors or for NIOSH to look at that and see if that approach

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appears to be -- would meet their needs. mean we've -- in other petitions we've had, you know, NIOSH saying they haven't done it yet but they can do it, and then we said we'd like to see some examples. But it seems to me there may -- they may be far enough along in their development of some of these that if NIOSH took a look at that -- it wouldn't yet be public -if that could be shared I think that might be an interesting -- or a possibility to -- to see whether or not the -- I mean NIOSH is saying they don't think such an approach is possible, so you know, when these come out, then it either is possible or it isn't, so I wonder if some kind of discussion with those, if -- if that was possible, it wouldn't have to be -wait till it's peer-reviewed and published. Some part of a peer review would be NIOSH taking a look to see what they're doing and if the justification for how their interim have been going are consistent with a more rigorous science they may bring to bear might be a -one thing that could be done quite quickly. DR. ZIEMER: I think at the moment you're not necessarily proposing that, but asking for

1 perhaps a reaction from --2 DR. ANDERSON: (Off microphone) 3 (Unintelligible) saying it can't be shared --4 DR. ZIEMER: -- from the two agencies --5 DR. ANDERSON: -- (unintelligible) it's out of the --6 7 DR. ZIEMER: -- and -- and while we're letting 8 them ponder that a moment, Rich, did you have 9 an additional follow-up on that? 10 MR. ESPINOSA: No, not necessarily on that. I 11 was just -- I'm trying to wonder in my mind, 12 what were the living conditions? Did these --13 were these workers shipped to the islands and 14 then taken off or did -- were there -- living 15 at the island? I'm trying to think of like 16 time-weighted averages. 17 DR. ZIEMER: Uh-huh. I'm not -- I don't know 18 if either -- either Paul or Jim can answer 19 that, or the petitioner, whether -- whether Ms. 20 Karo can answer that or knows, but --21 DR. NETON: I really can't -- can't answer that 22 question. The problem with these cases is very 23 much like a lot of our cases where we -- we 24 lack a lot of information. We have a case 25 file. We have a -- potentially a job title,

1 and we know the person wore an external badge. 2 We know how long they were there, and that's about it. So unlike --3 4 DR. ZIEMER: Whether they were back and forth, 5 you don't really know. DR. NETON: Yeah, and some of that is very hard 6 7 to figure out. On top of that, unlike the 8 military records, which tend to be a little 9 better as far as where these people were 10 stationed, we have statements that say the 11 person was there, and we don't know where they 12 were and we kind of know in general what they 13 may have done. So we have access to far less 14 information about what these people were doing 15 and where they were than even the military may 16 have. 17 Jim, while you're at the mike can DR. ZIEMER: 18 you react to Dr. Anderson's question? 19 I suppose. I think --DR. NETON: 20 DR. ZIEMER: Or maybe Larry Elliott will react. 21 DR. NETON: Maybe I'd better let Larry react to 22 that. 23 DR. ZIEMER: Let the boss react to that. 24 MR. ELLIOTT: No, I don't want to answer, but I 25 will try to answer. I think it certainly is a

possi--

MR. ELLIOTT: It is a possibility. I was just conferring with Dr. Blake, and I think we need to -- it's a possibility that -- that could be pursued, but it's going to result in time expended. And we're okay with that if that's the pleasure of the Board. I think -- you know, my concern that I expressed to Dr. Blake was what kind of perturbation would our entrance into your review and approval process present to you, and they have an advisory body, as well. So my answer is yes, it's possible; it comes with consequences and some of those consequences we may not be able to fully characterize at this point in time.

DR. ZIEMER: Well, we're just exploring, is

DR. ZIEMER: Uh-huh.

DR. WADE: Dr. Blake, you mentioned you would be presenting to your advisory board. When would that be?

DR. BLAKE: The interactions I have with my advisory board are basically two-fold. On all these technical publications that we're pushing ahead, they are looking at them now. I have

not asked for a -- we have actual sub-groups on our advisory board, and the sub-group that interacts and does technical reviews -- in my board there's a subcommittee number one on dose reconstruction that's simply focused on our processes. The chairman is a member of the -- one of the members of the NRC report from 2003. They're providing a peer-review process of the publications that are generated by my agency through some of our contractors, too. But I will --

The second part of it is when we actually have formal Veterans Advisory Boards. Our next one is due in June of this year, or July, in Austin, Texas. And at that time I will be -- as Dr. Neton mentioned, we released a report to Congress from both the Department of Veterans Affairs and Department of Defense, mandated by a Public Law, and I need to get back with them on our get-well plan formally at that time where we are. And so the majority of my formal comments at that time will be in June.

DR. WADE: Thank you.

DR. ZIEMER: Jim Melius.

DR. MELIUS: Yeah, I -- two comments. First

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addressing Richard's question, I think it was a very good point and -- about where did -- where were these people living when they were at the site and where they stayed. That was actually one of the rationales for the -- Amchitka not having the 250-day requirement was the people doing the work there lived on that island and there were questions about their exposures sort of 24 hours a day and sort of how do you weight that and so forth. So I think that should be pursued, and I would also wonder whether there wouldn't be information -- probably not in the interviews 'cause I don't think that's asked for -- but some follow-up. I don't know to what extent the claimants are still alive, how many there are, if any, or -- but some of that kind of issue could be pursued and information sought from the survivors or from other -other information that -- that might be available, so I think that deserves follow-up. I think back to Henry's comment and really sort of -- sort of question, I think -- understand, you know, the reason that yeah, there's some initial time limits, but -- and there's also an issue that we may not be able to totally

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reconcile the way that dose reconstructions are done in these two programs because they have different legislative mandates and follow different methods. However, to the -- I would think we need to be very careful about sort of institutionalizing a difference between the programs and -- and the way that they're approaching or utilizing certain types of data, simply 'cause we didn't want to, you know, wait a few months for, you know, a report to get finished or -- or do a complete evaluation of that. And we would be making a decision on a Special Exposure Cohort that would -- is going to go forward for a long period of time, and yet it may turn out that in a few months or several months that some of these issues that we used as the basis -- might use as the basis for approving a Special Exposure Cohort may be addressed, and therefore we would have institutionalized a difference that wasn't really part of the difference in the legislative mandates for these two programs. And I think there's a need for -- I would -even though there may be differences in the way people worked and there are differences in the

laws, I think we need to be -- try to maintain as much consistency as we can, and certainly not have sort of unnecessary inconsistency between the two programs. So I would certainly feel much more comfortable waiting until we could try to clarify some of these technical issues.

DR. ZIEMER: Thank you. Roy DeHart.

DR. DEHART: Again this issue of disparity between populations. We're sitting in Oak Ridge, and I can assure you there is an enormous emotional difference between people who are awarded at X-10 and who are awarded at K-25. That difference is painful for many of these people and what we -- it sounds like we're potentially going to be doing, there may be grounds to do it, creating two different systems for same kinds of exposure settings, and I think we need to move slowly there.

DR. ZIEMER: Rich, I think your -- Rich
Espinosa's comment is one that it would be nice
if that could be followed up on. It seems to
me that it's fairly obvious many of these
people were indeed in place 24 hours a day on
those atolls and so on and therefore maybe one

would consider looking at the 250-day issue in a somewhat different way for this kind of a situation. Larry Elliott.

MR. ELLIOTT: Thank you. I just want to make two comments and -- just for the Board's consideration and understanding. At least 57 claims that you see on the screen that remain to be dealt with, some of those come in in the first batch we received in October of 2001. Given that, you know, I'd just like to put that on the record for those claimants who are awaiting a decision.

Secondly, I would note in the three that have been treated with dose reconstructions, one of which that was found to be compensable had multiple skin cancers and we did that using external dose. The other two were found to be non-compensable and I don't know the exact cancer type, but I think both of those were cancers that are in the presumptive list. So just offer that for your information.

DR. ZIEMER: Michael.

MR. GIBSON: Yeah, on page 31 of 38 of the petition in the book, the one dated 10/20, I believe it is, at the top of the page it talks

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about (reading) considerable attention to recreational activity, personnel in the Entwok (sic) island had two movie theaters, a TV station, a hobby shop, a swimming pool, beach areas designated for swimming, a skeet range, playing fields, basketball and handball court -- handball courts and a service club with a snack bar, library, game room, rooms for adult education classes and clubs. Competitive leagues were organized for many sports. Number one, it would take quite a length of time to construct such facilities, I would thing. And number two, I don't think that they would go to that trouble, it wouldn't appear to me, unless there was some extended amount of time that people would have to stay on the islands.

DR. ZIEMER: A good point, Michael. I want to ask -- maybe I'll ask Paul Blake if he can answer this, but -- or maybe Jim Neton can. We know that in some cases -- for example, the Marshall Islands where there was a shift in the wind and the fallout really was concentrated on one of the islands; they actually evacuated those folks and they essentially have been off

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the island ever since. It's been decades. These locations were ones where the -- I don't believe -- they were not in the direct cloud or they wouldn't have had the people stationed there. They must have gotten more peripheral fallout, I presume. Do we have some idea of -there were some air samples, were there not, that give us some idea of the levels, and you're using some kind of resuspension factor so you must know something about these levels that people were living with every day. And we were, incidentally, in this country, as well. We did take extensive radioactive DR. BLAKE: fallout measurements there. But there were cases, as I mentioned earlier, where the wind did shift and there were some unexpected consequences and that, you know, was challenging to us. But most of that data not only comes from film badge, but it's also from the actual measured fallout -- some of which was on land, but some of it was in -- obviously in water samples, biological samples. was one group that was mentioned in Dr. Neton's review, the University of Washington, that has been doing this work for many years. So we

have a number of different sources of data that
we use in actually calculating this, but
radioactive fallout was measured fairly
effectively.

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DR. MELIUS: Yeah, I think -- goes back to my -- my first question I had about how people qualified as -- issue of being badged, or should have been badged. And I guess my concern also would be that we may not be qualifying people in the correct way for a Special Exposure Cohort that -- it's not saying that one might not be justified for some of these simply based on the amount of information available on their activities and -- and taking into account where they lived, some of -- some of the other uncer-- certainties here, but I just think we would be able to do a much better job and much fairer job for everybody involved if we had more information to be able to work from on -- on this and in -- particularly some of -- either the additional scientific work that's underway for the veterans, as well as some further information, some further work on the part of NIOSH in sort of describing this group of people in this -- this cohort and sort

of figure out how we -- how we best address the different groups that there are there.

DR. ZIEMER: Wanda Munn.

MS. MUNN: I don't know that what I'm thinking adds anything to the debate, but there's no question that there's -- in my mind, that there's a difference between military personnel and civilian personnel. There's a difference in the way they were assigned and the way they are monitored -- were monitored.

If the real question here is did people do the same work, one set being military and the other being civilian, and are those individuals with the same exposures being treated differently, I submit that it is unlikely we are ever going to be able to have that information clearly in hand at this stage. It was too far removed from the actual events.

If it is also true that for most individual who were long-term on the island the primary driver is likely to be internal exposure, then our choice appears to be to accept the fact that we will never be able to assess parallel activities between military and civilian personnel in that group, and accept the fact

that we will be unlikely at this moment to evaluate what internal dosimetry -- excuse me, what internal doses were experienced by the civilian workers who were there.

If we cannot do that, then our civilian workers were in -- are in a definitely different category than the military workers whose exposure was characterized as being more external than internal.

I don't know how long we can postpone making the decision, and whether we can be very sure that the DTRA process is going to be applicable for us. If -- if we delay based on the assumption that it will be and it's not, then I suppose what we have to be reconciled to is the fact that we will have to propose to those people who are -- have been waiting for four years for us to move forward that we're not ready to move forward yet, we're still working on it. That seems a very difficult choice, for me, and I don't see any real assurance that we'll be a lot better off once we see what Dr. Blake has been able to complete with his program.

Am I -- am I too far off, Dr. Blake?

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DR. ZIEMER: All the way.

DR. BLAKE: Not completely, but I would like to mention -- during the Pacific Proving Ground exercises the -- and detonations there, it was a joint task force that worked together of both the AEC personnel and DOD personnel. For instance, external dosimetry was -- used the same types for both. And in fact the -- the records now for the most part are maintained -for the military, it's jointly funded by both DOE and my program out in Las Vegas where we have many of the older film badges that were used. We've kept actual physical hard copies, plus documentation of the film badges. Doses are actually maintained out at the Nuclear Test Archive in Las Vegas. It's a DOE-run facility that I jointly, as program manager, co-fund. So during all those years, at least with external dosimetry, we tried very much to have consistent methodologies between both AEC and the military. Internal dose, though, does remain a challenge, and there was no good solution really at that time.

MS. MUNN: And --

DR. ZIEMER: Mr. Presley has some related

comments.

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MR. PRESLEY: As one that did not work on an above-ground shot, but one that has worked on quite a few below-ground shots, you would have a large contingency of what we would call at that time (on microphone) AEC or ERDA personnel that would be the group of people that would be handling not only the shot material but the shot equipment and things like this. have DOD there. They were -- and they had the Seabees and things like that, as Roy mentioned. They were more of an observer. At that time and present today, your AEC/DOE still handled the everyday business of working with the material, putting the weapons together, getting it up in the air or down in the hole, whatever it was. We did that as a -- as an atomic worker contractor.

Then we would go back in and do the sampling after the shots, and I have no reason to think that things were any different, especially from 1956 to '62 when they were working on 17, 35 and 36 shots at a time. You would have a tremendous people (sic) getting ready for each one of these shots. You didn't just go in in a

1 day and pop one and walk out and go back the 2 next day and pop another one. It didn't happen 3 that way. 4 So yes, they were -- there was a lot of people 5 there. They did have permanent party facilities. I would think that there's 6 7 probably about the same thing we used to have 8 at NTS in the early days was you lived in 9 quonset huts or tents and it was -- there would 10 be quite a few people working on that kind of 11 stuff. I -- I would not think that it would be any different -- might be a whole lot more 12 13 modern, but it would be any different from when 14 I started out there in the '70s than it was 15 back in the '50s or the '60s. 16 DR. ZIEMER: But you're suggesting then that 17 the -- the military portion would be more 18 transient than the -- the folks doing the shots 19 are actually the AEC ERTA/ERDA folks. 20 That's correct. MR. PRESLEY: 21 DR. BLAKE: If I could comment, from Operation 22 CROSSROADS on, even though it was a joint task 23 force, there were really two functions. 24 AEC people were focused on the weapon. And the 25 military was there primarily for weapons

effects. We needed to understand those effects and how our troops are going to move ahead.

And so it was a different function with the two groups. But in the Pacific Proving Grounds many of those tests were weapons effects tests and we brought in large amounts of military personnel because of that.

DR. ZIEMER: Yeah. Thank you. Okay, Gen Roessler and Jim Melius.

DR. ROESSLER: I feel like Wanda, I'm not sure what I can add to this. But I do want to say what I'm thinking at this point. I'm pretty much convinced that we're dealing with two different populations, for the most part, and that would persuade me to go ahead and vote for this petition at this time.

However, I've been sitting here listening to the comments from everybody around the table and from Dr. Blake and Dr. Neton, and I've been going back and forth with my vote. I feel very uncomfortable with that. I feel like maybe I don't have enough information at this point in time to -- to vote, even though if right now I voted, it would be for the petition. And I -- I wouldn't be concerned that much about the

1 difference in the agency approaches because I 2 think it is a difference in the type of 3 exposures and type of population. 4 DR. ZIEMER: Thank you. Jim Melius. 5 DR. MELIUS: Jim, were you going to answer? DR. NETON: Well, I was going to comment on 6 7 something a couple --8 DR. ZIEMER: Oh, sure. 9 DR. NETON: -- of persons ago, so if I could --10 DR. ZIEMER: Jim Neton, go ahead. 11 DR. NETON: -- interject here just briefly. 12 I'd like to follow up on the issue of the joint 13 task force and badging. It's true what Dr. 14 Blake said that there were joint task forces 15 and the badging was handled by -- and 16 consolidated in Nevada Test Site. In fact, of 17 the 57 people that I show we have external 18 badge results for, that's where we obtained the 19 results, the film badge results resided in the 20 Nevada Test Site. 21 The difference I see is that these people, even 22 though they were on the joint task force and 23 were badged, we have very little information 24 about what they were doing and where they were 25 the entire time. It's also complicated by the

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fact that from 1958, I believe, the REDWING series, not everybody was badged. They started badging people I think around -- after '58, and prior to '58 it was what we have come to call in this group "cohort badging". You know, certain people were badged. So then we faced the dilemma of having to reconstruct external dose to civilians based not -- knowing of their whereabouts and using environmental survey results to reconstruct the external, to then turn around and reconstruct the internal based on the reconstructed external. It just multiplies on top of each other. So it is a somewhat more difficult process, I think, than what you would see with the military. DR. MELIUS: Yeah, actually I think it -- were sort of two issues that I think need to be

addressed, but I think the main point is that DTRA -- Dr. Blake says that they have some significant, you know, technical information currently being developed that would address the main technical basis for the SEC determination, the issue of internal monitoring. And as I understand it, they've offered to -- now to make that information

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available to NIOSH and -- in some way through -- through a process. And I would think that it would be appropriate for us to delay consideration of this petition till our next meeting. In the meantime, ask NIOSH to address two things. One is work with DTRA and, you know, take a look at this technical information and make a determination, is it going to address the concerns or not, and sort of -- you know, what is -- what would be the timing for that, how -- how long would it take before this information could or could not be used for dose reconstructions -- is it going to be adequate (unintelligible) that. And I -- again, for the reasons I've said before, I -- I'm just uncomfortable -- you know, if in a few month suddenly our concerns have been addressed. Secondly, I do think we need to also address this 250-day issue. With the people living there -- again, it may not be significant exposure because of location or whatever, it may be able to be taken into account in the dose reconstruction, but it is something different and I think we need to be careful for this circumstance in terms of how we qualify

1 these -- these people. So I would ask NIOSH to 2 also address that over the next -- between --3 by our next meeting, at least to --4 DR. ZIEMER: I'm --5 DR. MELIUS: -- come back with some additional information to us on --6 7 DR. ZIEMER: I'm not going to recognize that at 8 this point as a motion, but as a possible 9 motion. I want to hear from Roy DeHart and 10 then Richard Espinosa. 11 UNIDENTIFIED: (Off microphone) 12 (Unintelligible) 13 DR. ZIEMER: Oh, okay. 14 DR. DEHART: I've just been signaled that I'm I propose the following motion: 15 That the 16 Board thank NIOSH for its report related to the 17 SEC petition; further that the Board move to 18 table further discussion to a time certain to 19 allow possible conferences and discussion 20 between NIOSH and DTRA and to address other 21 issues as previously stated by the Board. 22 This in essence is a motion to DR. ZIEMER: 23 table. Is there a second? 24 MR. PRESLEY: I second. 25 DR. ZIEMER: This is not a debatable motion.

1 We must immediately vote. This is a vote to 2 table, and it's to a time certain, which is the 3 next face-to-face meeting. 4 Just by way of instruction, if you do not wish 5 to table -- and there could be an alternate 6 motion, such as Jim suggested. If you do not 7 wish to table, which ends our debate today, 8 then you will vote no. If you wish to table 9 with a time certain that it comes off the table 10 at the next meeting, you will vote yes. 11 Any procedural questions? 12 (No responses) 13 Okay, we're going to vote by hand -- raising 14 your hand. All in favor of tabling the action 15 -- motion to table any action on this petition 16 -- actually the Chair's going to change the 17 ruling. We have nothing -- we do not have a 18 motion before us to table. Roy, we don't have 19 a motion before us to table. 20 UNIDENTIFIED: (Off microphone) 21 (Unintelligible) don't have a motion 22 (unintelligible). 23 DR. MELIUS: There's nothing to table. 24 DR. ZIEMER: There's nothing to table, 25 actually.

1 DR. DEHART: Okay. 2 DR. ZIEMER: What we would -- what we would 3 need would be --4 MR. PRESLEY: A motion --5 DR. ZIEMER: -- a proposed action, which could then be tabled. 6 7 DR. DEHART: Yes. 8 DR. ZIEMER: We have -- we have a petition 9 evaluation report, which in essence comes as a 10 request for action. I guess -- we still need a 11 motion to -- to do something with that, so --12 but hang on, we'll get a comment from Jim. 13 DR. NETON: I'd just like to make one comment. 14 I want the Board to understand the reality of 15 what's going to happen if this thing gets 16 pushed to the next Board meeting, which is --17 in discussion with Dr. Blake, it's unlikely 18 that we will -- NIOSH will receive any peer-19 reviewed report in time to be -- to evaluate 20 any changes on our decisions here. So it may 21 quite be likely that I would come back here in 22 April and say we -- we have not seen anything 23 fleshed out in detail. 24 Now there may be some preliminary approach

design documents that he could provide to us,

1 but there is unlikely to be -- in time for the 2 next Board meeting, us to have additional 3 information to weigh in on. 4 DR. ZIEMER: Thank you. Mike? 5 MR. GIBSON: It seems to me that we as a Board 6 have been very concerned about time limits, the 7 180-day time limit, dragging things on and 8 doing things in a timely manner, and I just 9 wonder if tabling this motion would buy us 10 anything. 11 DR. ZIEMER: Thank you. We're not actually 12 discussing a motion to table 'cause it's not discussable (sic), but you're talking 13 14 theoretically, if one were to make such a 15 motion -- okay. 16 Robert Presley. 17 MR. PRESLEY: I'd like to make a motion that we 18 accept this petition, as read. 19 DR. ZIEMER: Okay, that's a valid motion. 20 there a second? 21 MS. MUNN: Second. 22 DR. ZIEMER: Now we have before us an action. 23 That is a motion to accept the petition -- and 24 actually it's to accept the recommendation of 25 NIOSH to grant the SEC petition, and if this

1 motion passed it would be a recommendation to 2 the Secretary of Health and Human Services that 3 SEC status be granted. This is a debatable 4 motion. 5 Wanda. I have one question for NIOSH, or 6 MS. MUNN: 7 for Dr. Blake, whoever is best qualified to 8 answer it. Can we make the assumption, or do 9 we know for a fact, that resuspension and 10 internal dosimetry for individuals on those 11 islands would vary from location to location on 12 the islands? Can we make that statement? 13 DR. BLAKE: Yes, they certainly do vary. 14 MS. MUNN: They vary. 15 Yes, they do vary. DR. BLAKE: 16 **UNIDENTIFIED:** Can you hear that, Ray? 17 MS. MUNN: If they vary, then I am back to my 18 comment earlier. I do not believe it is 19 possible for us to ever define where these workers were at any given time on that island 20 21 and therefore cannot, with any degree of 22 specificity, come to a conclusion with respect 23 to their internal dose. 24 DR. ZIEMER: So you are speaking for the 25 motion.

1 MS. MUNN: I am speaking for the motion. 2 DR. ZIEMER: Jim Melius. 3 DR. MELIUS: Yeah, I would just like to point 4 out on Wanda's point that that is not one of 5 the claims that -- one of the claims that NIOSH has made in its evaluation. I don't believe 6 7 that point was raised, unless I --8 DR. NETON: That -- the resuspension factor is 9 one of the key issues raised by the National 10 Research Council. 11 DR. MELIUS: Okay. 12 DR. NETON: And the uncertainty associated with 13 the resuspension values --14 DR. MELIUS: Is it uncertainty of location or 15 uncertainty --16 DR. NETON: Not on location, but uncertainty of 17 -- of -- well, they were talking about uncertainty of resuspension on the ships versus 18 19 the island and such, but there is significant 20 uncertainty associated with the resuspension 21 factor. 22 DR. MELIUS: I was more reminding us that if we 23 were going to use that for a basis we needed to 24 -- for an SEC, we needed to capture that 25 specifically in however we make our

1 recommendation.

My question, though, is back -- I thought I understood Dr. Blake to say that he was going to make some of the technical work that was currently underway available to NIOSH prior to it being published in peer review journals, and I --

DR. WADE: Dr. Blake --

DR. MELIUS: -- the sort of discrepancy here in terms of -- of what Jim and Larry just said and what I understood you to say earlier, I'm just trying to understand it and --

DR. WADE: Could you please clarify, sir, just what would happen between now and the end of April in terms of interactions between your staff and NIOSH toward shedding light on this issue of internal exposure?

DR. BLAKE: When I get back tomorrow to my own agency, I can release some of our preliminary design documents on how we're actually moving towards these publications. Many of them are basically written, but they need to go through a formal peer review process, not -- at least for the DTRA technical reports, before we publish them and put them perhaps in a slightly

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more readable format. Then our plans are to actually put them through our subcommittee 1 dose reconstruction board to have an outside look at them before we move ahead. The report of greatest concern right now is one that's due out -- that I release some preliminary information on tomorrow, but it's only about a two-page summary. The report itself is an update on the FEDOS, the software code model. That's not due for publication until about the end of March or April. And then with a peer review process, let's assuming a month in putting it in, it won't be ready for your next Board. I can't promise that. And that's the reason I said end of calendar year 2006. are a number of other technical reports that are -- that may be a little more advanced than that. But once again, I do have only limited number of personnel to get them out in a timely manner.

DR. ZIEMER: Paul, are you able -- well, number one, you're saying you have no guarantee that the end product will look like what you have now since there's some iterations that you have to go through.

1	DR. BLAKE: True.
2	DR. ZIEMER: Number two, are you able to
3	personally commit your agency to sharing this
4	information outright with another agency at
5	this point? Are you authorized to make that
6	commitment? Well
7	DR. BLAKE: I believe I believe I am, sir,
8	and I don't
9	DR. ZIEMER: Well, I just want to make sure.
10	DR. BLAKE: I don't expect that'll be a
11	problem, but whenever you ask someone in the
12	federal government
13	DR. ZIEMER: Well
14	DR. BLAKE: are you authorized, we always
15	take a breath before we
16	DR. ZIEMER: as far as you know.
17	DR. BLAKE: As far as I know, sir, yes.
18	DR. ZIEMER: Roy DeHart and then Henry and
19	Rich.
20	DR. DEHART: I propose the motion that you have
21	previously heard as to tabling.
22	DR. ZIEMER: Okay. We now have a motion to
23	table this motion, and that proposed motion to
24	table was to table to our next meeting. And
25	was there a second to the motion to table?

1	DR. MELIUS: I'll second that.
2	DR. ZIEMER: Okay, this is a motion to table
3	the action that was proposed by Mr. Presley.
4	All in favor of tabling, raise your right hand.
5	One, two, three, four, five.
6	Those opposed?
7	One, two, three, four
8	MR. ESPINOSA: Abstain.
9	DR. ZIEMER: One abstention. We have what,
10	we have five
11	DR. WADE: Five to four with one abstention.
12	DR. ZIEMER: four and one. The Chair voted.
13	We have ten. Tabling requires a majority of
14	those voting, but the abstention is not
15	ignored. We have it still requires six, I
16	believe, is unless I get overruled,
17	Parliamentarian-wise.
18	MR. PRESLEY: Could I change my vote?
19	DR. ZIEMER: You want to change your vote?
20	Yes, you can.
21	MR. PRESLEY: I'll vote to table.
22	DR. ZIEMER: We now have six votes
23	DR. WADE: Let's take a let's take the vote
24	again, please.
25	DR. ZIEMER: Raise your right hand if you're

1 voting to table. 2 One, two, three, four, five, six. Okay. 3 THE COURT REPORTER: Dr. Ziemer, who changed 4 their vote? Was that --5 DR. ZIEMER: Mr. Presley. The vote is to table 6 the action until the next meeting, and it is so 7 ordered. 8 Now I'm going to have a brief recess, and after 9 the recess I'm going to ask for a motion that 10 will provide instruction to NIOSH as to what 11 should occur between now and the next meeting, 12 since the motion to table leaves things in a vacuum. Okay? And we'll get the additional 13 14 comments at that point. 15 Let us recess for 15 minutes, and then 16 reconvene. Okay, Daniella, if you're still on 17 the line, you may wish to remain. We will try 18 to provide some instruction to NIOSH on -- as 19 to what should occur. 20 MS. KARO: Thank you, sir. 21 (Whereupon, a recess was taken from 10:30 a.m. 22 to 10:55 a.m.) 23 DR. ZIEMER: Okay, let us proceed. We do have 24 some additional action to take. Having tabled 25 the action on the Pacific Proving Grounds, we

1 do need to provide some instruction as to what 2 we expect to happen within the next week. 3 Also, in terms of information from another 4 agency, we do have some advice from legal 5 counsel. I'm searching for the right word, but Liz, can you advise us in terms of what we can 6 7 and can't do with information from another 8 agency? 9 MS. HOMOKI-TITUS: Certainly. Liz Homoki-10 Titus, (unintelligible) HHS. I'm not sure it's 11 advice, it's just a concern that you need to be 12 aware of that if another department provides 13 information to HHS that is not publicly 14 available -- I guess I just want to expand on 15 Dr. Ziemer's question to DOD regarding the 16 release of that information to the Advisory 17 Board and to the public. Obviously when 18 information is released to a FACA advisory 19 board -- can you hear me okay? 20 UNIDENTIFIED: (Off microphone) 21 (Unintelligible) and start over. MS. HOMOKI-TITUS: (Off microphone) You want me 22 23 to start again? 24 THE COURT REPORTER: Excuse me? 25 DR. ZIEMER: Ray, does she need to start again?

1 THE COURT REPORTER: Oh, no, you can go 2 forward. Thanks. 3 MS. HOMOKI-TITUS: Okay. When it's released to 4 a FACA advisory board it's obviously made 5 public, or we would have to take the Board into 6 closed session to consider that information, so 7 I just want you to be aware of those possible 8 options. And I don't know if the 9 representative from DOD wants to address that 10 concern --11 DR. ZIEMER: Well --MS. HOMOKI-TITUS: -- regarding that 12 information --13 14 DR. ZIEMER: -- at least be aware --15 MS. HOMOKI-TITUS: -- becoming public. 16 DR. ZIEMER: -- aware of that, and it may be 17 that if, for example, we suggest that NIOSH get 18 together with DTRA on that issue, that Jim 19 Neton or his designee can report back to us 20 whether they found they could utilize whatever 21 is being done, without necessarily releasing 22 information at that point would be another 23 option. But I guess we do want to make sure 24 Dr. Blake is aware that if the information 25 fully comes to the Board, it gets made public.

1 So that's -- that's the point. 2 Now the Chair will recognize Dr. Melius for 3 purposes of making a motion. 4 DR. MELIUS: Yeah, I would like to -- not only 5 would I like to, but I am making a motion --6 DR. ZIEMER: He doesn't like to, but he will do 7 it. 8 DR. MELIUS: -- to make a motion to request 9 that NIOSH follow up on three items in 10 relationship to this petition. 11 Number one was what Liz was just referring to -12 - to, which is to gather further information, 13 evaluate the current technical work that's 14 underway at DTRA, that -- particularly that 15 addressing the internal dose monitoring and 16 evaluation of internal dose. 17 Number two, we're asking NIOSH to conduct 18 further evaluation of the types of work and the 19 work pattern for people working at the Pacific 20 Proving Ground, particularly we're trying to 21 better understand what types of work, what the 22 nature of the exposures for people in different 23 work categories, and also addressing the 24 clarification on -- to what extent people were

living there and people who were in residence

1 there, how they were exposed as part of their 2 living condition. 3 And number three, specifically how to address this -- best to address this issue of 4 5 qualification for the cohort, both the classification as having -- being monitored or 6 7 likely to -- I think it's supposed to be 8 monitored or should have been monitored, in the 9 context of these workers. And also whether 10 it's appropriate to it if people were residing 11 at the site and exposed during their 12 residential hours, should the 250-day 13 requirement be adjusted to take that -- that 14 into account for those workers. Remember, we 15 have a 250-day work-day requirement and that 16 may not quite make sense in terms of -- it 17 doesn't make sense in terms of people are --18 are living on the site and exposed, you know, 19 24 hours a day, essentially -- at least in some 20 manner, and I think that --DR. ZIEMER: Okay. 21 22 DR. MELIUS: -- that needs to be --23 DR. ZIEMER: Right. 24 DR. MELIUS: -- addressed in the context of the

1 DR. ZIEMER: Now you can --2 DR. MELIUS: -- Special Exposure Cohort. 3 DR. ZIEMER: -- speak in support of the motion 4 in a moment, if it's seconded. 5 DR. MELIUS: Okay. DR. ZIEMER: Is there a second to the motion? 6 7 MR. ESPINOSA: I second. 8 DR. ZIEMER: Now it's seconded. 9 debate, pro or con, for this motion? 10 speaking in favor of the motion? 11 THE COURT REPORTER: Dr. Ziemer, who seconded? 12 MS. MUNN: DeHart. 13 DR. ZIEMER: Okay. DeHart seconded. DeHart? 14 Jim, I'd like to ask you to clarify for the 15 Chair, when you said address the qualification 16 of should have been monitored, what are you 17 asking for there? Is that a different request 18 than the evaluation of the 250-day requirement? 19 DR. MELIUS: I think it's part of that. - it's slightly different, but in light of the 20 21 different groups that are being evaluated or 22 could be potentially included in a Special 23 Exposure Cohort, I just wanted to make sure 24 that that particular requirement makes sense

for -- is appropriate for different groups.

1 And I think we're going to find that there were 2 a number of different categories of workers 3 here and I just want to make sure that's 4 appropriate for those -- those workers. Again 5 we go back to (unintelligible) what does it mean to be monitored or should have been 6 7 monitored. 8 DR. ZIEMER: Do we know whether -- where that 9 monitoring occurred for people stationed -- was 10 that monitoring 24 hours a day as opposed to a 11 usual worker that leaves their badge at the 12 gate when they leave a facility? Paul, do you know the -- whether that was the case or not? 13 14 Or did these workers wear their badges --15 DR. BLAKE: (Off microphone) I don't know 16 (unintelligible), sir. 17 DR. ZIEMER: Okay. Paul says he doesn't know. 18 Robert? 19 MR. PRESLEY: When I worked at the test site 20 you wore your badge everywhere you went. You 21 were on the site all the time, 24 hours a day, 22 7 days a week. 23 DR. ZIEMER: Okay. Thank you, that's helpful. 24 I think Rich Espinosa and then Michael. 25 MR. ESPINOSA: I speak in favor of the motion.

My biggest concern is the -- under the law, the 250 days. I believe that if it was passed today, that's -- might -- the way the law reads that a lot of these people that are defined under the class would be denied on the basis of 250 days, so I speak in favor of the motion and I do believe that this Board could look for ways and get around and make a recommendation.

DR. ZIEMER: Thank you. Michael.

MR. GIBSON: I also speak in favor of the motion and I would just offer perhaps a friendly amendment. If there's not issues dealing with classified information or if there is, if there's a member of the Board that's got the proper clearance, could a member of the Board be present during these discussions with NIOSH and DTRA?

DR. ZIEMER: I think the answer is yes, that could be done.

MS. HOMOKI-TITUS: It's not a question of clearance, it's simply that these are internal documents that haven't been released to the public, so yes, a working group of the Board could work with NIOSH in those meetings. Can you hear me okay? But it's a matter of when it

1	becomes public if it's given to all of the
2	Board.
3	DR. ZIEMER: Thank you. Other questions or
4	comments?
5	(No responses)
6	Are you ready to vote on the motion then?
7	MR. GRIFFON: Was that accepted as a friendly
8	amendment?
9	DR. MELIUS: Yes.
10	MR. GRIFFON: Okay.
11	DR. MELIUS: Very friendly.
12	DR. ZIEMER: So point one of the motion asking
13	asking NIOSH to gather further information
14	from DTRA on internal dose would include the
15	caveat that this be done with a Board member or
16	members present.
17	Are you ready to vote on the motion?
18	UNIDENTIFIED: Yes.
19	DR. ZIEMER: All then in favor, say aye?
20	(Affirmative responses)
21	Those opposed, no?
22	(No responses)
23	Any abstentions? One abstention, Wanda Munn
24	abstaining.
25	Motion carries. Thank you very much. I don't

1	know if Daniella's still on the line.
2	MS. KARO: Yes, actually I am.
3	DR. ZIEMER: Do you have any additional
4	comments
5	MS. KARO: No, not really. Not at this point.
6	DR. ZIEMER: Then we will make sure that we
7	make arrangements to have you present at the
8	next meeting, by phone if necessary, and
9	perhaps we can arrange the schedule so that it
10	is not quite so early in the morning.
11	MS. KARO: Yes, it would be greatly
12	appreciated.
13	DR. ZIEMER: Okay. And we thank you for
14	participating with us today.
15	MS. KARO: Thank you.
16	DR. ZIEMER: You're welcome to stay on, but you
17	probably would rather do something else.
18	MS. KARO: That's true.
19	DR. ZIEMER: Wouldn't we all.
20	DR. WADE: Not me. CLASSIFIED DATA: IMPACT ON BOARD SEC PETITION RECOMMENDATION DR. LEWIS WADE, EXECUTIVE SECRETARY
21	DR. ZIEMER: Okay. Next on our agenda we have
22	an item called classified data, impact on Board
23	SEC petition recommendation, and Dr. Wade has
24	that issue. And I believe we have a document -

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DR. WADE: I've given out a --

DR. ZIEMER: -- that's been distributed called
"Classified Information".

DR. WADE: And it's on the table. Before I begin that, I would like to, for the record, point out that I'm operating on the belief that following this meeting the new members will be duly impaneled. So if you were to start to consider Board members listening in or participating, then we have the new members to consider.

Okay, to the issue on the agenda. I think you'll all remember this issue. It was at one point a very emotional issue. On your behalf - you had written to the Secretary of HHS. On your behalf I met with representatives of the Secretary's office and representatives of HHS's Office of General Counsel last week to -- to try and bring this issue to a head. Based upon those discussions they empowered me to come to you with this position. So the position I bring you is the HHS position, and let me walk you through it.

Under classified information, the first bullet

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says (reading) Based upon legal advice of the Department of Justice, Office of Legal Counsel, it has been concluded that non-disclosure to the public of classified or restricted information does not qualify a class for addition to the SEC if a sufficiently accurate dose reconstruction is otherwise feasible using classified or restricted information.

I pointedly asked the people I met with if this opinion from the Department of Justice was rendered in writing. I was told that it was not. It was a verbal opinion of the Department of Justice that formed the basis of this written statement that HHS is making. I asked them again, and they assured me that there is no written opinion of the Department of Justice.

The second bullet says (reading) Therefore, the Secretary -- read HHS -- has no legal authority to grant a Special Exposure Cohort petition because classified or restricted information was used to determine that a sufficiently accurate dose reconstruction can be done. Again, that bullet relates to the Secretary and

the way the Secretary must act. We'll talk at

1 the end about my recommendation as to how you 2 should act. 3 And then the third bullet, (reading) The 4 Department of Justice has also indicated that 5 access by claimants or the public at large to classified or restricted information on which 6 7 HHS may rely in making its feasibility 8 determination is not required by due process 9 considerations. 10 Item: Petitioners have the opportunity for an 11 administrative review within the Department (as 12 provided by the SEC Final Rule). 13 And Item: If the petitioner files a lawsuit 14 and the court concludes it is necessary, the court can review the classified information ex-15 16 parte, in camera. 17 So again, you had asked about the due process 18 considerations. Now, again, this isn't new 19 information I'm bringing to you, but this is 20 information that last week I went to the 21 Secretary's representatives and HHS Office of 22 Legal Counsel, and this is the information they 23 asked me to bring to you. 24 The only thing I would add, on the second 25 bullet, it speaks to the Secretary's

1 responsibility. As your Designated Federal 2 Official, I think it is most appropriate that 3 you be mindful of the Secretary's 4 responsibility in your framing of 5 recommendations to the Secretary. I in no way limit your ability to say what you think needs 6 7 to be said. I do think you need to be mindful 8 of the Secretary's responsibility, again, as 9 you frame recommendations to the Secretary. 10 DR. ZIEMER: And let me ask, Board members, do 11 you have any questions for Lew on this issue? 12 Michael. 13 MR. GIBSON: If that's the guidance from the 14 Office of Legal Counsel, what's the resistance 15 to putting that in writing so that it can be 16 made public and -- to the Board and to the 17 public so -- in an effort to be transparent? 18 DR. ZIEMER: This --19 DR. WADE: I think this --20 DR. ZIEMER: -- this is public. 21 DR. WADE: This is public. 22 MR. GIBSON: But that's just the HHS opinion of 23 what they --24 DR. ZIEMER: Right. 25 MR. GIBSON: -- they determined the Office of

1 Legal Counsel said. 2 DR. WADE: Right. 3 MR. GIBSON: I'm wondering why the Office of 4 Legal Counsel is resistant to put it in writing 5 for us to look at. 6 DR. WADE: Well, I don't think it -- I can't 7 answer that, Mike, though I think the opinion 8 that's relevant is the opinion of the Secretary 9 and the Office of Legal Counsel of HHS, and 10 they're willing to put their recommendation in 11 writing. They based that in part on 12 discussions with the Department of Justice, and 13 they do not have that in writing. I can go 14 back and ask them again to try and get it in 15 writing. It's the wish of the Board with 16 regard to that. They do not have it in writing 17 at this point. 18 My principal concern is the Board be not 19 limited in its ability to do its business, and 20 this is what I wanted to be able to bring to 21 you. 22 DR. ZIEMER: Jim Melius. 23 DR. MELIUS: Yeah, I mean I would argue that 24 this does limit the Board in its ability to do 25 its work -- essentially disenfranchises us in

circumstances where there is classified or restricted information that, you know, many or -- if not all members of the Board would not have -- have access to and would preclude us from essentially evaluating NIOSH's review of the SEC petition. We would not be able to, you know, perform any meaningful review of the -- of that -- of that petition. So I would argue that it doesn't nec-- it certainly doesn't facilitate the Board doing anything. In fact, it -- it restricts and, in essence, could disenfranchise us in certain -- certain situations.

I'd also remind us that we do have I believe a letter that has never been answered, other than this recent verbal transmission to you, Lew, requesting that written opinion. I mean -- and for reasons that in order to -- for the Board to be able to operate. I think we had good reasons then. I just I believe summarized some of them that I -- trying to remember back there, there could have -- there were more -- additional ones, but I don't think it's a very workable approach, given the nature of this program. And particularly it certainly

undermines any transparency for this program and any credibility that the Board adds to the recommendations. And are we supposed to just, you know, defer any review of a NIOSH recommen- recommendation on an SEC simply 'cause we won't be able to technically review it or do an adequate review of it in circumstances -- and those circumstances, do -- what happens, does it just pass through? Do we write a letter saying sorry, we can't do anything?

DR. ZIEMER: Liz?

MS. HOMOKI-TITUS: I would just remind the Board that EEOICPA actually gives the Department of Energy authority to provide Q clearances for Board members. So I don't think this limits Board members' access to classified information. This limits public access to classified information.

DR. WADE: And we do have Board members who have clearances. It could be that you wish to increase the number, and then we could work towards that end. The -- the model we've been working on is that there are Board members with Q clearances that would be privy to this information and then could report in summary to

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the Board. If you feel there needs to be a greater number of Board members so cleared, then we could pursue that.

DR. MELIUS: That's certainly one element of it, but it would certainly change how the Board would review and (unintelligible) and I guess, given some of the conflicts of interest, it -it's -- it would require a large number of people. We have some significant problems, for example, on Y-12, by the way, as to who would be -- if anybody would be available to review the -- review classified information. Do that.

DR. WADE: I understand.

DR. ZIEMER: We actually have had some difficulty in getting people Q cleared in the past. I know that it's a pretty long, tedious process and -- is there -- is there any -- one of the -- one of the problems is, unless you do it in advance -- if you wait till you have a petition in hand and say we need O-cleared people, it -- it -- the time factor is very long. How practical is it to think about getting a large number of this Board Q-cleared? DR. WADE: We could start now. It's not reasonable to assume that it would be done in

1 any short term. 2 DR. MELIUS: And can I also just add that even 3 with Q clearance there's access on like, what, a need-to-know basis and -- and there are still 4 5 -- information that's still restricted. clearance isn't automatic access to all --6 7 DR. ZIEMER: Right. 8 DR. MELIUS: -- the classified information, and 9 there are situations where, if I recall 10 correctly, that NIOSH staff is not allowed 11 access to the --12 MR. ELLIOTT: No, that's not true. 13 DR. MELIUS: -- information. 14 MR. ELLIOTT: We -- the need-to-know has been 15 established and it's covered in the memorandum 16 of understanding between HHS and DOE. 17 our experience in getting staff and contractor 18 staff clearances depends on an individual set 19 of circumstances for each case. It runs from 20 nine months to a year. 21 DR. MELIUS: Uh-huh. 22 MR. ELLIOTT: It can be expedited if you've had 23 a previous clearance and, in some cases, that's still taken six months' time. 24

I'd also note for you, though, that while we

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have finished dose reconstructions for 12,000 claims, not one of those have used classified information to arrive at a dose reconstruction for decision. But this -- this counsel that's been provided would also cover those cases or claims where we might have to use classified information to adjudicate a final decision. I just want to make that point clear. only SEC petitions that are at risk here, it's also perhaps a claim or two in the future. DR. MELIUS: Can I ask a question related to that point? I'm not -- I don't know the answer and it may be something Liz or Larry -- do your procedures currently -- your regulations currently address this -- that part of the issue, the individual claimants' access to this in-- this information? 'Cause if I recall, we set up a process and the law sets up a process, I think presuming that there is -- there is access, and -- and certainly that transparency was an important element of it, and does that MR. ELLIOTT: The law does -- the law does speak to the use of classified information. EEOICPA recognizes that there is a need for

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national security regarding certain types of information, and that we will end up facing Our processes, our rules, do not speak that. to how an individual claimant or a petitioner may be afforded access or may not be afforded access. We have sought guidance and counsel from our own general counsel and from the Department of Labor's general counsel, and the Department of Justice general counsel. We have been told that there are procedures that are employed by the Department of Justice to address how and whether or not access can be given. And you know, I -- to date we've not had to gain experience in any of those procedures so I can't speak to how cumbersome or how difficult or how successful they may in fact be.

DR. MELIUS: Uh-huh.

DR. WADE: And just for -- for information on the table, SC&A also has Q-cleared people that are available to the Board.

DR. ZIEMER: Further comments? Michael.

MR. GIBSON: Based on this, just to throw this out for a thought for the Board, should we draft a letter to Congress and indicate to them

1 that due to this latest information, it may 2 preclude us from fulfilling our obligations 3 fully? 4 DR. ZIEMER: I'm not sure who the question is 5 addressed to, maybe to ourselves. I suspect 6 that if we wanted to raise that issue, we would 7 probably want to do it through the Secretary as 8 a concern, either asking the Secretary to do 9 something on our behalf or take the route of 10 requesting that Board members be granted Q 11 clearance. Lew, do you --12 DR. WADE: No, I agree. I mean I think that's 13 the normal way to proceed. It -- it's really 14 up to you, but... 15 MR. GIBSON: Based on the fact we already have 16 the opinion of HHS, could I make a motion that 17 we draft a letter to the members of Congress 18 and inform them that, based on this 19 information, it's -- may preclude us from 20 fulfilling our obligation. 21 DR. ZIEMER: Your motion is that we draft a 22 letter to Congress? 23 MR. GIBSON: Yes. 24 DR. ZIEMER: Okay. Is there a second to the 25 motion?

1 (No responses) 2 Is there a second? No second? 3 DR. MELIUS: Well, I'm just asking a procedural 4 question 'cause I thought either Lew or Paul 5 indicated it would be a letter --6 DR. ZIEMER: Well, I was suggesting that --DR. MELIUS: -- to the Secretary. 7 8 DR. ZIEMER: -- I -- I thought any such letter 9 would go through the Secretary. But that does 10 not bind the Board. If the Board wishes to 11 directly correspond with Congress, I think we 12 can make such an action. We do correspond with 13 Congressional people on a regular basis. It's 14 getting more and more regular, it seems. 15 DR. WADE: I do think this is an advisory board 16 to the Secretary. 17 DR. ZIEMER: I'm not hearing a second yet. not sure if that's discomfort with --18 19 DR. MELIUS: It's -- well, personally it's I'm 20 not sure the best way is to go to Congress. 21 certainly would propose that we communicate with the Secretary -- two-fold, reiterate our 22 23 request for the written opinion from Department 24 of Justice 'cause I think -- I'd like to 25 understand the rationale and sort of -- better

1 understanding how we may -- how we 2 operationalize this very general sort of 3 statement from -- from them that's, you know --4 at least how HHS interprets something verbally 5 said by some verbal ruling from the Department 6 of Justice, and I just find it... 7 DR. ZIEMER: Let me make a suggestion, just to 8 move us forward on this issue. I'm going to 9 suggest, Michael, that you and Jim get together 10 and draft a proposed motion for the Board for 11 us to act on tomorrow on this issue. 12 we understand the intent is to elevate this 13 item and make the consequences known. Perhaps 14 you and Jim could -- is it -- if that's 15 agreeable -- could prepare a written motion for 16 us to act on at our work session tomorrow. 17 Would that be agreeable? 18 DR. MELIUS: That'd be fine. And it should --19 should that motion suggest a letter, we'll 20 draft a letter, also. 21 DR. ZIEMER: That would be fine. 22 DR. MELIUS: That way I think will 23 facilitate... 24 DR. ZIEMER: That would facilitate. 25 DR. MELIUS: Okay, good.

DR. ZIEMER: Okay, we'll proceed on that basis, then we don't have to wordsmith something right here.

BOARD CORRESPONDENCE RESPONSE TO LETTERS FROM
SEN. CLINTON, SEN. SCHUMER, REP. HIGGINS, REP.
SLAUGHTER, AND MIKE WRIGHT, STEEL WORKERS
(PRESENTATION BY NIOSH/ORAU ON WORKER INTERACTIONS)
DR. PAUL ZIEMER, CHAIR

Well, speaking of Congressional correspondence, you have in your notebook several letters. And under the Board rules, the Chair has drafted some responses. I'd like to begin with the letter from Senator Clinton, which -- a letter was dated November 7th dealing with Bethlehem Steel. A proposed response was drafted and distributed to the Board earlier this month by e-mail. Copies of it are on the table. The e-mail transmission dropped the -- big word "draft" should be on this. This is not a final letter, it's a draft. It's not been sent. The draft is dated January 19th. It's in your notebook.

This is the proposed response to Senator
Clinton's letter. I would like to point out
that I also sent a draft of this to Stu
Hinnefeld at NIOSH and asked him to provide me
with updated figures on the Bethlehem Steel
site as far as -- because the draft letter to

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Senator Clinton used the -- I think it was the November data from the site, so I did ask for that update, and somewhere I have that information. Now it's not in the draft copy that you have. I have a -- it'll just take me a moment here to make files out of it. I'm going to ask for a motion to accept or to transmit my proposed letter, with the following changes. In paragraph two where it says "Department of Labor has referred 678 cases", that number now is 692, from Bethlehem Steel for dose reconstruction. Dose reconstructions have been completed on 578 cases -- and you'll notice that number has dropped from the original value of 580, and that's because a couple of cases apparently got referred back or something like that, so the correct number now is 578, and that is -- instead of 85 percent, it's 83 percent of the total cases, so make that change. The last sentence in that second paragraph should now say three cases have been withdrawn by DOL, leaving 111 cases at NIOSH awaiting dose reconstruction. The next paragraph would say of the 578, 255 -that number remains the same -- appear to have

1 a probability of causation of 50 percent or 2 greater, and 323 will have a probability of 3 causation less than 50 percent. According to 4 Department of Labor, compensation paid to 5 Bethlehem Steel workers as of January 19th, 2006 was \$38,250,000, with an additional 6 7 \$194,139 in medical bills paid. 8 So those are the updated figures, and then the rest of the letter would remain the same. 9 10 like to ask for a motion to transmit this 11 letter. 12 DR. WADE: Just a minute. You are -- we are 13 also in receipt, about the same time frame, 14 from -- letters from Congresswoman Slaughter 15 and Congressman Higgins --16 DR. ZIEMER: Right, and those were --17 DR. WADE: -- and you were --18 DR. ZIEMER: -- those are separate letters, 19 yeah, and we'll handle them individually since 20 the letters are different. 21 MR. PRESLEY: I'd make a motion we accept this 22 letter to be sent. 23 DR. ZIEMER: Is there a second? 24 MS. MUNN: Second. 25 DR. ZIEMER: Now I'd call for any modifications

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that anyone wishes to make or suggest -- Jim.

DR. MELIUS: Yeah. The draft that's in front of me, it's the last paragraph on the first page, (reading) In light of the actions already taken by the Board...

DR. ZIEMER: Uh-huh.

DR. MELIUS: I -- I object to this way this paragraph is written. I understand the rationale, but it seems to me that Senator Clinton's letter makes the assumption that the Board generates a Special Exposure Cohort, and then we sort of say well, we're not planning on doing that. Well, we can't do it. It's not -you know, we act on NIOSH's rec-recommendations. NIOSH either, because they can't complete dose reconstructions in some cases, are forwarding SEC to us, or you know, the more common case so far has been with, you know, petition and people petition us to do that. So I mean I think we need to, you know, address that directly and say it's not -- you know, not our task. We're not in a position to -- you know, we have not been asked to consider an SEC petition there. And I'm a little concerned about -- I think we can say that a

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number have been done, but we're in this very awkward position. We review individual dose reconstructions. We have I believe reviewed some from Bethlehem. But we may have -- I can't remember if we deferred them or what the action was, but -- but it's only a small number. We have a site profile that we've reviewed and what -- you know, the -- you know, it's still in process. I mean NIOSH is revising the site profile. And there's sort of a lot of -- there's sort of a lot of process, things that have been done but haven't been finalized here. So I think we need to be careful in reaching conclusions about something we're not in a position to reach conclusions about because procedurally we haven't done it. I mean I understand what you're trying to say there.

DR. ZIEMER: Well, actually I thought I was trying to say just that and perhaps didn't say it well. Number one, the action already taken is referring to the paragraph where we gave an opinion. It's quoted.

DR. MELIUS: Uh-huh.

DR. ZIEMER: That's the action, that -- that

the site profile is adequate for dose reconstruction. That's a key point, I thought.

DR. MELIUS: Yeah.

DR. ZIEMER: Number two, that there is no petition. I mean the last paragraph says there is no qualified petition on which the Board could act. Maybe if you feel that Senator Clinton doesn't understand the process, I'd -- we can certainly expand that. I was simply pointing out, in essence, that the Board cannot do anything -- we cannot act, there's -- for SEC, there's no petition before us to act on.

DR. MELIUS: I would --

DR. ZIEMER: So help us say it better.

DR. MELIUS: Yeah, I would bring that to the front of the paragraph 'cause what I'm concerned about, there are no plans by the Board to recommend a special exposure cohort -- I mean where we don't plan unless we're asked to review. That -- that -- that's what -- DR. ZIEMER: I gotcha, okay. So suppose we start the paragraph with -- we drop the word "furthermore" and say "At the present time there's no qualified Bethlehem Steel exposure cohort on which to act."

1 DR. MELIUS: Right, yeah. 2 DR. ZIEMER: Then what? In light of actions 3 already taken and because the -- and then just 4 drop the thing -- no plans for the Board to 5 recommend -- since we don't do that, that's the 6 point that you are making, I guess. 7 DR. MELIUS: Yeah. 8 MR. ESPINOSA: Drop that whole sentence? 9 DR. ZIEMER: I think the suggestion is that we 10 drop it, starting with -- let's see -- well, 11 I'm not sure -- we can't just drop that phrase, 12 though. 13 DR. MELIUS: I think our official position is 14 better said in the paragraph (unintelligible), the one that starts "On January 9th". 15 16 we've stated that we voted on that and we -- I 17 think we really have sort of formally done what is says here. And because NIOSH --18 19 DR. ZIEMER: Well, I --20 DR. MELIUS: -- has demonstrated... 21 DR. ZIEMER: When the Board instructed me on 22 January 9th, the instruction was to deal with 23 her request for -- that we act on -- or declare 24 an SEC petition, so we do need to say that at

the present time there's no qualified --

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1	DR. MELIUS: Yeah, I think I think that
2	and I would
3	DR. ZIEMER: And then just end at that point.
4	DR. MELIUS: I think just end at that point.
5	DR. ZIEMER: Is everybody comfortable with
6	that?
7	MR. GRIFFON: What what's happening with
8	that whole sentence before, in "In light
9	of", I think that's
10	DR. ZIEMER: That would disappear.
11	MR. GRIFFON: Then I'm then I'm satisfied.
12	DR. ZIEMER: I certainly consider that
13	friendly. It's within the idea
14	DR. MELIUS: Yeah.
15	MR. GRIFFON: Yeah.
16	DR. ZIEMER: Wanda, did you have a comment or
17	suggestion? I found no dangling participles.
18	MS. MUNN: I'll be cautious about that. Do I
19	understand correctly that you're proposing to
20	eliminate the entire first sentence of the last
21	paragraph?
22	DR. ZIEMER: Yeah, it's the second to last
23	paragraph and it would simply say "At the
24	present time there's no
25	MS. MUNN: Yes.

1	DR. ZIEMER: "qualified Bethlehem Steel SEC
2	petition on which the Board can act."
3	MS. MUNN: If our purpose is to help instruct
4	the Senator in process here, then would it not
5	be wise to say "At the present time there is no
6	Bethlehem Steel Special Exposure Cohort
7	petition that has been qualified by the federal
8	agencies and on which the Board could act."
9	Would that not be clearer? Or would it simply
10	muddy the waters further for staff? You know
11	staff better than I, Dr. Melius.
12	DR. ZIEMER: I can add that (unintelligible)
13	simply it might imply that there was a
14	petition but it didn't qualify. I don't think
15	there's been any petition. Any other
16	suggestions?
17	(No responses)
18	Then with that change, let me ask for a vote on
19	this draft.
20	All in favor, aye?
21	(Affirmative responses)
22	Opposed?
23	(No responses)
24	The motion carries and we will transmit that
25	letter.

1	Then we have a memo let me take the Schumer
2	letter next. This also deals with Bethlehem
3	Steel. Oh, incidentally, this letter would be
4	accompanied by the matrix that I referred to in
5	the letter.
6	MR. GRIFFON: Yeah, attached, you said.
7	DR. ZIEMER: Yeah, it would be attached. We
8	don't have that here as part of it, but
9	Now the Schumer letter that's in your file is
10	not the one to which I was responding.
11	DR. WADE: There are two.
12	DR. ZIEMER: Oh, there's two, okay.
13	DR. WADE: And November 14th is the one to
14	which you were responding.
15	DR. ZIEMER: November 14th, and then I'm
16	looking for my response, here it is.
17	DR. WADE: Your response.
18	DR. ZIEMER: Okay, here's the response, and
19	there is a similar paragraph in the Schumer
20	letter similar to the Clinton letter that in
21	fact, I believe it's identical. But let me ask
22	for a motion to transmit this letter, and then
23	we can take care of it.
24	MR. PRESLEY: So moved.
25	DR. ZIEMER: Second? Moved by Presley,

1	seconded by
2	DR. DEHART: Second.
3	DR. ZIEMER: DeHart. Does the Board wish me
4	to modify that second to last paragraph in the
5	same manner as the Clinton letter?
6	DR. MELIUS: In a very friendly way, we would
7	ask you to do that.
8	DR. ZIEMER: Any other any other amendments
9	or modifications?
10	(No responses)
11	If not
12	MR. GRIFFON: Is there is there any update
13	on on Linde in here? I'm just reading this
14	quickly now, but the the letter was asking
15	about Linde Ceramics workers as well as
16	Bethlehem Steel.
17	DR. ZIEMER: Well, he was thanking us for our
18	action on Linde.
19	MR. GRIFFON: Right, okay, okay right. I'm
20	just refreshing myself, sorry before I vote.
21	DR. ZIEMER: Okay, all in favor, aye?
22	(Affirmative responses)
23	Opposed?
24	(No responses)
25	The motion carries. Now we have

1 DR. WADE: The second Schumer letter. 2 DR. ZIEMER: -- the second Schumer letter, 3 which is dated --4 **DR. WADE:** Coincidentally, January 19th. 5 is new to the Board, there's a second Schumer 6 letter dated January 19th that has recently come to Dr. Ziemer. It has an attachment, a 7 8 two-page document that has Eddie Walker's name 9 at the bottom of it. That should be in your 10 (unintelligible). 11 MR. PRESLEY: That's two pages? 12 DR. WADE: Right, so you've got this letter 13 from Schumer dated the 19th, and then a two-14 page one from Eddie Walker that leads with "Dear Mr. Elliott". 15 16 DR. ZIEMER: What is the date of the Ed Walker 17 -- is that the same one we got 18 (unintelligible)? 19 MS. MUNN: Yes, I believe it is. 20 DR. MELIUS: My recollection is when we 21 considered Bethlehem on the January 9th call 22 that we discussed dealing with Mr. Walker's 23 concerns and that we -- that our action there, 24 in terms of instructions to NIOSH on resolving 25 and changing the site profile, presumed that,

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should there be further correspondence or issues raised by Mr. Walker, that NIOSH would address those in, you know, an on-- sort of an ongoing manner in terms of, you know, similar they'd addressed to other issues raised for other site profiles, you know, and for claimants and would take those into account in an appropriate fashion. Is that -- am -- I'm -- probably asking -- I thought that was my recollection 'cause we talked about sort of how to address Mr. Walker's concerns. And not all of them were addressed in -- in our instructions in terms of the site profile and the issues we were trying to resolve. And -and I would suggest that we write back to Senator Schumer something to that effect, that it's our understanding that -- and that NIOSH's actions, you know, in -- revised site profile at some point is going to come back to the Board. NIOSH is going to provide us quarterly reports, et cetera.

MR. ELLIOTT: That is correct, as I recall, as well. And we have been responding to the numerous letters that Mr. Walker has provided us. This recent letter, we'll again respond to

1 the points that he raises, as we have, and we 2 will post these on our web site, as we do, so -3 - and we intend fully to bring forward to the 4 Board whatever we resolve in that matrix of I 5 think six or seven items, so... DR. WADE: Is Mr. Walker either in the room or 6 7 on the telephone? 8 DR. ZIEMER: He's not in here at the meeting. 9 DR. WADE: And he's not on the telephone? 10 Okay, I just want to make... 11 DR. ZIEMER: Let me suggest the following since 12 I had not seen the letter -- these letters 13 often, as this one did, don't come to me 14 directly so there's a time lag. I had not seen 15 the letter, but I -- the Board's policy is that 16 -- that letters to Congressional people obtain 17 Board approval. I'm going to suggest that I draft a letter along the lines of what was just 18 19 described and distribute it to the Board by e-20 mail and see if there's any major objections 21 with the wording. If there is not, we'll just 22 go ahead and transmit it. If there is, we'll 23 delay sending it. Is that agreeable? 24 MS. MUNN: Please do, please. 25 DR. ZIEMER: Any objection?

1 (No responses) 2 Without objection, we'll do that. 3 DR. WADE: So just so it's understood, the 4 procedure -- Paul will draft a letter, send it 5 to you. If any Board member raises an 6 objection, then we'll hold for the next 7 meeting. If no Board member raises an 8 objection, then Paul will be empowered to send 9 the letter. 10 DR. ZIEMER: It'll simply provide a response 11 along the lines of what you described, Jim, 12 that NIOSH is going to bring to us any changes 13 in the site profile. 14 DR. MELIUS: Right. 15 DR. ZIEMER: He's asking here that we rescind 16 the action we took on January 9th. And unless 17 I hear action to that --18 DR. MELIUS: No, I -- 'cause --19 DR. ZIEMER: -- I'm going to tell him that we 20 are not rescinding that action. 21 DR. MELIUS: Yeah. No, I -- and because our 22 action took into account that further 23 information may be raised and --24 DR. DEHART: (Off microphone) I don't 25 understand what his concern was.

1 (unintelligible) we didn't take action. 2 DR. MELIUS: And I think --3 DR. ZIEMER: No, we acted on that motion on 4 January 9th, and I can point out to him that 5 that motion did take into consideration the findings of SC&A. He implies here that we 6 7 should rescind that because of SC&A's findings, 8 and we took those into consideration. 9 DR. MELIUS: But our action didn't tell NIOSH 10 not to -- to ignore any further --11 DR. ZIEMER: No. 12 DR. MELIUS: -- correspondence or --13 DR. ZIEMER: That's right. 14 DR. MELIUS: -- information that came in from 15 Mr. Walker or anybody else, and -- and I think 16 that's -- that's sort of the point. It's not 17 like we've -- that's all that'll ever be 18 considered are those six points, but rather 19 than as new information becomes available it 20 would be incorporated. 21 DR. DEHART: That was my recollection. DR. MELIUS: Yeah. No, I'm -- 'cause I asked 22 23 the question specifically. I was trying to 24 understand the -- that, because Mr. Walker with 25 -- some of the information was not sent in a --

1 you know, in a -- at least in a timely way that 2 could be addressed by our resolution process and -- and so forth. 3 4 DR. ZIEMER: Okay. Thank you. Now there was 5 another --DR. WADE: Two other Congressional --6 7 DR. ZIEMER: -- series of letters. 8 DR. WADE: These are letters from Congresswoman 9 Slaughter and Congressman Higgins. My 10 recollection was that -- it was a feeling that 11 the Clinton letter would suffice in response, 12 but you need to make that determination. 13 DR. ZIEMER: The Louise Slaughter letter --14 wasn't this a statement to the Board at the 15 time of our meeting, as opposed -- this was not 16 a letter to me. Wasn't this a public statement 17 made by her -- by her staffer? DR. WADE: It's possible. 18 19 DR. ZIEMER: My recollection is that this --20 this Slaughter statement was read into the 21 record already and -- well, it's just part of 22 the deliberations, as was the Higgins -- these 23 are just written copies of what was read into 24 the record, so the Board officially received 25 those. I don't think any action is necessary

1 on those. 2 DR. WADE: That's fine. 3 DR. ZIEMER: There was one additional letter 4 that asked for certain information, and -- and 5 the Board has been copied on this. I'm -- I 6 don't see it in our booklet --7 DR. WADE: Who was it from, Paul? 8 This was -- this was the --DR. ZIEMER: 9 actually we will get to it, it's a Hanford 10 letter. 11 DR. WADE: Right, that's --12 DR. ZIEMER: We'll cover that when we get to 13 Hanford. 14 DR. WADE: Right. 15 DR. ZIEMER: And the information was provided 16 directly I think, Lew, by you or by NIOSH and 17 we'll cover that at the appropriate time. 18 DR. WADE: Right. 19 DR. ZIEMER: Okay. 20 DR. WADE: The only other longstanding letter 21 we have is a letter from Mike Wright with the 22 steelworkers. We've arranged for ORAU to brief 23 the Board on its efforts with regard to 24 collecting and taking into account worker 25 information. We could have that briefing now

1 or we could have it after lunch, as you 2 would... 3 DR. ZIEMER: How long does that briefing take? 4 MS. KIMPAN: I intend to be very brief, 5 depending on the questions and comments the 6 folks may have. 7 DR. ZIEMER: Why don't we go ahead and do it, 8 we have about 15 minutes, don't we? Yes, let's 9 proceed. 10 DR. WADE: Right, the only thing I would ask --11 please, proceed, come to the microphone. I'd 12 like to hold to the 1:30 time slot for the 13 beginning of the Rocky Flats discussions again 14 'cause we'll have folks from Colorado on the 15 phone, but I think we're -- that's well within 16 our -- our ability to do. 17 DR. ZIEMER: Okay, so we'll proceed with this -18 19 MS. KIMPAN: (Unintelligible) Hello, thank you 20 Dr. Ziemer, Dr. Wade, members of the Board. 21 I'm Kate Kimpan, the project director for the ORAU team for this effort. It's a pleasure to 22 23 be here today and share with you some of what 24 we're doing in both response to Mr. Wright's 25 letter to the Board and others last June, but

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also in response to the growing need that we as a team have felt and that our colleagues at NIOSH have felt and instructed us to deal with. And that is we, as a part of this entire process, get a great deal of input from a number of interested parties around the country. In particular we conduct worker outreach meetings where we try to assure that workers who are represented have representatives there to share with us important information that they may have. In Mr. Wright's letter in June he raises concerns about how we're dealing with the information that we gather, what we've done with it, and what our response is. And I wanted to start by saying I know you're familiar with some of the activities that have gone on, but our team has developed, working closely with OCAS, a database to capture these The name of that database, in our comments. world of acronyms, is the WISPR database, which is the Worker Input to Site Profile Revisions. This database has the capacity both to capture comments that were made, but as importantly, after those comments are captured, to assure

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that the commenter and public understand what our team's response was to those comments. I will say that in everything that we've done up until now -- I've been on board since November, so when I say everything, it might be a more limited everything. In the work that was done prior to that, and we're continuing, we at these meetings, where we're there intentionally to gather input from workers, always have the team leader for the site profile document there. There's also always a representative from NIOSH in the room. I want to assure you that every comment that we get is extremely important. But I don't want you to think that we go into these meetings and might walk away with some extremely important comment with a delayed action upon it. We're endeavoring to make very clear to folks that have commented what we've done. And in several instances, about a dozen or so, we have communicated back with the representatives of organized labor that were as part of the meeting to assure them that they know what -how their input is valued and what it's done. You've heard many, many discussions through the

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many meetings as to the different processes and procedures that have emerged. And those of you that have been watching this closely realize many of the things that we're doing, including everything from revisions in documents to development of the database to revisions in the conflict of interest policy, are issues that were first raised in these arenas. So I think our project's responsiveness to this comment -these -- these comments and concerns has been empirical and it's been boisterous. A place where we have absolute room to improve is to coordinate how we assure that folks that have made comments in these meetings, folks who did not wish to remain anonymous, understand what we've done in response to those comments. We have a procedure developed now that's -it's -- it's one of the ORAU procedures that oversee how we deal with these comments. endeavoring to comply immediately with those procedures. We've right now entered into our database, which has the capacity, as I said, to both have the comment and to have our response to it showing, the comments from -- more than 600 comments from eight of the largest meetings

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at the largest facilities. As I say, we're going to -- we intend to work very closely with OCAS in coming days and weeks as we continue to ramp this up to assure that the concerns that Mr. Wright outlines and that others may have -are you hearing us; when you hear us, what do you do with it -- we need to assure people that yes, we're hearing you. We're listening. have found interesting and essential information. You hear them in these meetings, people will identify records, identify processes that only those -- those folks who were in those facilities really know, so they're incredibly valuable. And a place where I think we have great room to improve is to assure that workers, their families and their representatives understand what that value is. More importantly, empirically understand how we've dealt with each one of the comments. If it's a comment that's important and already assimilated, it doesn't mean we shouldn't respond; it means we need to let those folks know thank you for your comment, here's what our action is and response. It's not thoroughly developed exactly how we'll

communicate with everyone because these arenas are quite broad. Obviously if it's a group of steelworkers that were part of the meeting, communicating back with that local is a very effective way to assure that folks know what's occurred. If it's an area of workers that were represented by others, or not represented at all, it creates more challenges for us, and we're endeavoring to determine what the best way is to assure people that they're being heard.

Putting things on the web is a great way for I've heard concerns in my -- my sidebars some. through this meeting that of course there are many, many of these workers who might come make a comment that aren't likely to be able to look on the internet for the response. So we're going to work closely with OCAS to determine how best to assure that the word is getting out to the people in a productive way so they understand how essential their input and contribution to these documents and programs are. As you know, all the many, many, many documents that we've worked through this system thus far, none of them would have made it to

1 where they are without the essential input from 2 those folks who are working in those 3 facilities, alerted us to processes, problems, 4 records and occurrences. It's an absolutely 5 inherent part of us doing our job well. We 6 have room to improve on assuring that workers 7 and their representatives and their families 8 understand that we're hearing them and what 9 they're doing, and I'd welcome any questions or 10 comments or suggestions that you all have to 11 assure we're doing this well. 12 DR. ZIEMER: Kate, could you identify for the record -- if you have it at your fingertips --13 14 MS. KIMPAN: Yes. 15 DR. ZIEMER: -- the ORAU procedure number --16 MS. KIMPAN: I do, and it's ORAU Procedure 17 0097, Revision 00. It's titled "Conduct of the Worker Outreach Program", and it was approved 18 19 on December 29th of this past year. 20 DR. ZIEMER: Thank you. And I want to 21 determine or ask John Mauro, do you know, John, 22 if that procedure's on our list -- updated list 23 of procedures to review? 24 DR. MAURO: I do not --25 DR. ZIEMER: It may be too new to -- but we --

1 DR. MAURO: -- (unintelligible). 2 DR. ZIEMER: -- want to make sure --3 DR. MAURO: (Off microphone) (Unintelligible) 4 list that. 5 MS. KIMPAN: Does yours -- pardon me. Ziemer, does your list include Procedure 0031, 6 7 Technical Basis Document Development Review and 8 Approval, which was also updated 9 (unintelligible) --10 DR. ZIEMER: Kathy has the list here and just 11 is checking it. We don't need to know that right now. I'm basically saying we want to 12 13 make sure that -- that we have a look at what's 14 being done. 15 The other question I'm going to ask -- you 16 know, when we have our public comment periods 17 here, and all of that information is captured of course in the transcripts, we have always 18 19 sort of assumed that the appropriate people 20 will pick up the information. If it's a 21 Department of Labor issue, that their reps will 22 follow it. If it's a NIOSH issue, that NIOSH 23 will follow it and so on. But I'm wondering if 24 anyone has thought about capturing -- do you

folks capture the appropriate remarks from our

1 transcripts to enter into the comment database? 2 MS. KIMPAN: The reason I looked at Larry is 3 there was much that had gone before. I -- I 4 have to say I don't know whether there's a 5 formal review of transcripts. I know that 6 after each meeting the government 7 representatives and the ORAU team 8 representatives certainly, with their own notes 9 and experiences, when something emerges here 10 you sort of see us all huddle, so there's an 11 immediate exchange of information. 12 DR. ZIEMER: Right, right, I'm sort of getting at -- and the Board doesn't really have a -- in 13 14 place a formal procedure to track what occurs, 15 but we are aware that there's a wealth of 16 information --17 MS. KIMPAN: Absolutely. 18 DR. ZIEMER: -- that also emerges from the 19 public comment periods, and we don't want to 20 lose that. So that perhaps we can -- if the 21 Board agrees with this, would -- and if NIOSH 22 doesn't object, to at least have some means of 23 sort of checking that --24 MS. KIMPAN: Absolutely.

DR. ZIEMER: -- maybe looking at the

1 transcripts and make sure that the appropriate 2 items have been captured, if -- if needed. 3 MS. KIMPAN: Absolutely, and I believe --4 absolutely. MR. ELLIOTT: Yeah, you caught me --5 6 MS. KIMPAN: Sorry. 7 MR. ELLIOTT: -- not sleeping, but thinking 8 about something else, so -- but yes, I think 9 before Kate's arrival -- you know, Dr. Toohey 10 always attended these meetings or had some 11 other ORAU person attend. There were folks at 12 ORAU who perused the transcripts and captured 13 in what was called a Top Hat database, which 14 was an ATSDR software package that was totally 15 unusable by everybody else in the world and we 16 couldn't give SC&A access to it, actually. 17 got that converted over now to this WISPR 18 database, and I think -- I think there's still 19 resident now in that, there's still some 20 comments that were collected --21 DR. ZIEMER: Good. 22 MR. ELLIOTT: -- from Board meetings through 23 the transcript mechanism. 24 I would just offer this, that -- you know, you

saw me jump up a couple of times last night --

1 DR. ZIEMER: Sure. 2 MR. ELLIOTT: -- and I pull people out in the 3 hallway and we talk to them and we try to get 4 down to the bottom of the facts --5 DR. ZIEMER: And we know that there's follow-up 6 occurring. I'm really asking about -- 'cause you're really formalizing the process so 7 8 something doesn't fall through the cracks. 9 MR. ELLIOTT: Right, and we want to make sure 10 that where there is essential information 11 that's relevant to doing dose reconstructions, 12 that has impact on dose reconstructions, we 13 capture that --14 DR. ZIEMER: Right. 15 MR. ELLIOTT: -- and we -- we tell people how 16 we're utilizing that information. But at the 17 same time I'll say this to you, Paul, that a 18 lot of what we hear, in the end is not truly 19 relevant --20 DR. ZIEMER: Sure. 21 MR. ELLIOTT: -- and needs to be couched in a 22 site profile. It may be couched elsewhere in a 23 Technical Basis Document, it may be couched 24 elsewhere in a Technical Information Bulletin,

and we're -- and Kate's rightfully commented on

where we stand to make some very I think
considerable improvement is in those areas
where things that we hear that we don't you
know, we just don't believe they're going to,
you know, make any difference at the end of the
day, we need to get back to those people and
explain why.
DR. ZIEMER: Yeah. Yeah. So
MR. GRIFFON: Can I just ask a follow-up on
DR. ZIEMER: Sure.
MR. GRIFFON: that? In the WISPR database,
do we have access to that? I haven't seen it.
I just don't know if SC&A has
MR. ELLIOTT: I don't think you do yet, but I
think the plans that work is underway to
give you access, to give SC&A access. I don't
think they've actually got into it yet, but
that's
MS. KIMPAN: (Off microphone) (Unintelligible)
MR. ELLIOTT: our intent.
MS. KIMPAN: (unintelligible)
MR. GRIFFON: Does that include
MS. KIMPAN: (unintelligible)
MR. GRIFFON: you mentioned in your

1 presentation 600 comments or so --2 MS. KIMPAN: Yes. 3 MR. GRIFFON: -- from these worker outreach --4 MS. KIMPAN: Yes. 5 MR. GRIFFON: -- meetings, so those are rolled into the same database? 6 7 MS. KIMPAN: They are. 8 MR. GRIFFON: Okay. 9 MR. ELLIOTT: And maybe to -- maybe to help 10 Kate out a little bit, you know, comments come 11 in a variety of mechanisms. Some comments are 12 -- we prefer written comments. And our policy 13 is written comments are replied to in writing, 14 and so that -- you know, if a person goes to 15 the trouble to put pen to paper, we feel that 16 we owe them the courtesy of responding in 17 writing, as well. And that's -- that's pretty 18 well established. 19 We get comments in worker outreach meetings 20 that are verbal. We capture those in the minutes. The minutes do go up on our web site. 21 22 What we need to do a better job of, as we noted 23 here before, is getting back to those folks and 24 letting them know hey, here's how we treated

You may not have seen it, you know, on the

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web site or you may not know that it got caught in this information bulletin, or you know, we appreciate your thoughts and your comments, but you know, for dose reconstruction purposes, you know, there's no real relevance there and this is why.

We get comments of course from this Board meeting. We also get comments as we travel around in town hall meetings that are separate from the outreach effort. A year -- two summers ago we put on workshops for dose reconstruction and how we go about doing that, and we had -- and we invited union stewards, safety reps from organized labor, advocates for groups of people at different sites where there wasn't organized labor existing. We pulled those people in. We're going to do that again this summer, as well. I just want to get that on the record and so that we can, you know, have people start thinking about if they want -- they have an interest in attending those workshops. That's another source we get input.

MR. GRIFFON: Can I just --

DR. ZIEMER: Good. Thank you. Mark.

MR. GRIFFON: -- have a question on the WISPR

1 database, is it -- does that have comment and 2 resolution? 3 MS. KIMPAN: It does. 4 MR. GRIFFON: (Off microphone) I think that 5 would be important (unintelligible). MS. KIMPAN: (Off microphone) It has what's 6 7 (unintelligible) --8 DR. ZIEMER: So you're really tracking each 9 item --10 MS. KIMPAN: It does, and that was part of --11 the other limit on Top Hat was it was a capture 12 without, you know, the live part of what did you do. You need to know that every one of 13 14 those comments has been in my shop triaged to 15 the proper manager for resolution. So we're 16 dealing very promptly, and our policy has been 17 -- although, as Larry points out, we have room 18 to improve how people understand that -- always 19 has been immediately following those meetings -20 - after the minutes are approved, because the 21 minutes must be approved by the participants --22 those meetings are thoroughly -- those -- those 23 minutes are thoroughly scoured, the issues are 24 pulled out and given to the proper manager and

the responses come back. And you see the

response emerging in -- in, you know, things as broad as some of the -- the way that we're doing our overall work, but we didn't say to that commenter you're why we developed a coworker ritual. But it's because of what we've learned at these meetings, as you know, that many of our OTIBs and many of the other procedures and processes are put in place, because of what we've learned. This database does allow for resolution. And as a matter of fact, it's essential that you have a resolution before you can close out an item.

DR. ZIEMER: Uh-huh.

DR. WADE: Just to collect some items from the long discussion we've just had, I'll take it upon myself to see that the SC&A task to review procedures is modified to include 0097 Rev. 00.

MS. KIMPAN: And Dr. Wade, also would you add to that the Revised Proc. 0031, which is the Technical Basis Development document, which has been revised to accommodate these other changes? It's part of our formal TBD process now.

DR. WADE: I'll do that. I'll ask that every effort is made to make the WISPR database

1 available to the Board and SC&A and that we e-2 mail the Board when that availability has been 3 granted. And then I would ask that -- that the 4 minutes (sic) from these meetings, starting 5 with this meeting, be reviewed thoroughly and, 6 as appropriate, entry into the WISPR database 7 be made of comments that are taken here. 8 MS. KIMPAN: Absolutely. 9 MR. GRIFFON: I would -- I would also say it 10 might be useful, after we get the WISPR 11 database, to have another forum where we can 12 talk to the worker outreach (unintelligible) --13 MS. KIMPAN: Absolutely, we welcome input on 14 how to --15 MR. GRIFFON: -- (unintelligible) it would be 16 useful to look through comments and resolution 17 first before we pursue any (unintelligible). 18 MS. KIMPAN: Absolutely. 19 DR. ZIEMER: Dr. Melius. 20 DR. MELIUS: I have two questions. The first 21 is that I believe a union representative last 22 night had pointed out that he had submitted 23 comments on one of the site profiles -- I 24 believe he said two years ago, but maybe I 25 picked up the dates wrong -- and he knew that

1 they had been received, but there had been no 2 response to that. And -- and --3 MS. KIMPAN: I might have been in a sidebar, do 4 you -- who -- who was --5 DR. MELIUS: It was a public (unintelligible) -6 7 MS. KIMPAN: No, no, I mean who was that? 8 DR. ZIEMER: Yes, in fact, I asked him if he 9 could confirm that he had transmitted the 10 information, and he said he had confirmation of 11 delivery or something. 12 DR. MELIUS: And I just was curious about a 13 response to that. 14 MR. ELLIOTT: It was Mr. Glenn Bell, and yes, 15 we did receive the information. And I -- I 16 don't know if Bill Tankersley can help me out 17 here, but what he submitted to us were maps of 18 the site and other information. 19 rightfully, ORAU takes information that's given 20 to it and makes sure it goes through a 21 classification officer because some of this information, while it may be so marked as non-22 23 confidential, before we put it up on our web 24 site, we need to make sure that we're not going

to put something up that's (unintelligible) --

1	DR. ZIEMER: Well, especially site plans
2	nowadays are very
3	MR. ELLIOTT: And I believe that's where it's
4	at
5	DR. ZIEMER: a concern.
6	MR. ELLIOTT: it's still being
7	classification review, I think.
8	UNIDENTIFIED: (Off microphone)
9	(Unintelligible) specifically about those
10	documents (unintelligible).
11	MR. ELLIOTT: And I hope I'm not stepping out
12	of bounds here, but I think the reason why is
13	there's some building names and numbers on
14	there and we have to be careful about that.
15	DR. ZIEMER: Yeah.
16	DR. MELIUS: Okay, but I guess one of my
17	point is one, is that first of all, this
18	has taken I believe two years, if we've
19	MR. ELLIOTT: I've talked to him. I've talked
20	to him and I've sent e-mails back to him
21	DR. MELIUS: Okay, okay, that
22	MR. ELLIOTT: so you know, it's I think
23	he's frustrated in the fact that we can't seem
24	to shake it loose.
25	DR. MELIUS: Okay. But I mean I think it also

addresses our earlier issue about classified information, so I mean this -- see it in another example, one of the frustrations involved and -- and potential problems in dealing with that and it's not always straightforward.

My second question is that in terms of scheduling these meetings and so forth, is it a policy that when there's any significant revision to a site profile that there is a meeting held to get input and, you know, review of that significant revision with the people at the site? I mean...

MS. KIMPAN: It certainly has been the case that at sites where we get a great deal of comment there's a great deal of revision. We often have visited that site multiple times. I apologize, I don't know if -- if there's a formal proc-- I think there is not a formal procedure that says if there's any change, we will conduct another meeting. At the time, as you know, that the outreach meetings were growing in the form that they were in, it was a -- it was a slightly less broad agenda. Now obviously we're very interested in input for

everything from dose reconstructions -- well beyond just those documents, so I don't think we always conduct another meeting after there's been a change.

We do communicate in writing, and I have 30, 40 examples in my packet, with the representatives of organized labor. For instance, the steelworker meetings, we always send things back to those locals when there's been a change because of their comments.

DR. MELIUS: I mean I -- 'cause I would suggest that that become part of your policy/procedure, do that. Now again, if there's a revision underway and you're -- you address that at a meeting, that's -- another meeting, that's fine.

MS. KIMPAN: Sure.

DR. MELIUS: I don't think it has to be done necessarily twice or whatever. But -- but certainly as part of that process at some point, what one -- I -- I think that would be good because -- I mean, you know, this has been late in coming about. It's taken a lot of effort and a lot of urging on the part of the Board to get this process in place.

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Secondly, there's an awful lot of frustration out there because these site profiles have been in place. They're being used for literally thousands of dose reconstructions without an opportunity for meaningful input until relatively recently. And I think the more that the -- well, you may disagree with me, Larry, but that's the -- certainly the -- my feeling and the belief of a lot of other people out at the -- at these sites, and -- and a sense that you don't take people's comments into account, or to a very small extent. And I think to address that I think you really need to beef up and formalize the process for when you will do that so at least people will have reassurances that significant changes will not be made without the opportunity for input.

MR. ELLIOTT: Let me just say that -- I'm sorry. Yes, we do go back and we attend to any requests for a revisit, and we've documented that effort. And again, this is what we've been saying all along here. We do have room for improvement on getting back to folks and telling them where their comments stand.

MS. KIMPAN: And Dr. Melius, I assure you those

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comments are taken extremely seriously. think we have absolute room to improve on how we assure people that's the case. But I want you to know that, as science teams, both OCAS and ORAU take extremely seriously these comments. And I really do have a lengthy list, although we're short on time, of the many, many, many changes in our actual operations for those thousands of dose reconstructions that have been in response -- direct response to these comments. The fact that you bring up that you don't -- you weren't aware of that, and importantly, the folks making the comments might not be, shows us where we have vast room for improvement. But I assure you we're taking those comments extremely seriously as a And if someone raises an issue at program. those meetings that has immediate impact, both OCAS and the team leader are in that room in real time, and they don't wait until something else to incorporate that important information. And it's why you see things like some of the TIBs that we've developed, some of the OTIBs and some of the other processes that we've developed are in direct response to this input

1 that we've received. 2 DR. ZIEMER: Thank you. Mark has a comment. 3 MR. GRIFFON: Just a follow-up to what you just 4 said. You said you had a list of these -- and 5 even if it's in draft form, I think that'd be 6 useful, and if you have a pri-- if you can get 7 a print-out of it for the Board --8 MS. KIMPAN: Of? 9 MR. GRIFFON: -- and for the public. You just 10 said you had a list of examples of where you 11 had comments that -- that resulted in changes. 12 MS. KIMPAN: I -- I do indeed have a list, but 13 it's sort of my scrawled, bulleted what-we've-14 dones. 15 Oh, it's not --MR. GRIFFON: 16 MS. KIMPAN: I'd be glad to formalize a --17 MR. GRIFFON: I think maybe --18 MS. KIMPAN: Yes, absolutely --19 DR. ZIEMER: Maybe in the future that would be 20 something to share, yeah. 21 MS. KIMPAN: Absolutely. 22 DR. ZIEMER: Thank you. 23 MR. GRIFFON: Especially if we're still going 24 to be waiting for the WISPR database. I mean I 25 think that will all be in the WISPR database,

1 so if we have access to that, that might... 2 MS. KIMPAN: Yeah, WISPR went into production -3 - full production on the 15th of January, so 4 it's relatively recent, and I know we're 5 endeavoring to ensure that y'all have access. 6 DR. ZIEMER: Okay. Kate, thank you very much 7 for that presentation, and we're encouraged by 8 the direction that things are going with the 9 new database, and we'll look forward to updates 10 as we proceed. Thank you. 11 MS. KIMPAN: Thank you very much. 12 DR. ZIEMER: We're going to break for lunch. 13 We need to be back promptly at 1:30 to discuss 14 the Rocky Flats site profile. (Whereupon, a recess was taken from 12:10 p.m. 15 16 to 1:35 p.m.) 17 DR. ZIEMER: I'd like to call the meeting back to order, please. 18 19 Before we begin our regular agenda items, I 20 just want to mention to the Board and to those 21 assembled here, relative to the United 22 Steelworkers letter that was being discussed in 23 terms of the capturing of information from 24 workers and so on that we discussed just before 25 the break, I will send a formal reply to that

letter. When we originally got it last summer, it didn't appear that it needed a reply, it was simply some information. And it closed with please contact us if you have questions, was the way it ended. But I will reply to it and summarize what is being done in a formal way by the contractor to track the comments of workers and then follow up on them, so I just wanted to let you know that I will reply to that letter, which is the Michael Wright letter.

ROCKY FLATS SITE PROFILE - DISCUSSION/

PLAN OF ACTION

DR. PAUL ZIEMER, CHAIR

Now we're ready to begin our discussion of the Rocky Flats site profile. I want to find out whether anyone from the Rocky Flats or from -- from the site is -- is on the telephone line.

Do we have anyone remote from -- from Colorado?

MR. DEMAIORI: Tony DeMaiori with the United Steelworkers.

DR. ZIEMER: Thank you. Let me ask you to repeat the name for our court reporter again.

MR. DEMAIORI: Tony DeMaiori.

DR. ZIEMER: You may want to spell that for him.

MR. DEMAIORI: D-e-Capital M-a-i-o-r-i.

DR. ZIEMER: Thank you very much. Is there

1	anyone else from your group there?
2	MR. HILLER: We also have staff of the Colorado
3	Congressional delegation. I'm David Hiller
4	from Senator Salazar's office.
5	DR. ZIEMER: Thank you. Anyone else?
6	MS. ALBERG: Jeanette Alberg with Senator
7	Allard's office.
8	DR. ZIEMER: Welcome.
9	MS. BOLLER: Carolyn Boller from Congressman
10	Udall's office.
11	DR. ZIEMER: Thank you.
12	MS. WARDER: Amy Warder with Congressman
13	Beauprez.
14	DR. ZIEMER: Very good, thank you very much.
15	DR. WADE: Now we will be discussing a matrix.
16	Do you have copies of the matrix?
17	UNIDENTIFIED: Yes.
18	DR. WADE: And please, if there's any if you
19	have any questions in terms of your ability to
20	hear, please raise them. We want you to be
21	able to participate as fully as possible.
22	DR. ZIEMER: Now our our session this
23	afternoon, for the Board members here, will be
24	somewhat redundant in that for some of the
25	Board members we had a presentation yesterday

dealing with Rocky Flats that was made to our subcommittee. And the Board's contractor, SC&A, which has done the site profile review and represented here by Joe Fitzgerald, did present some material to the Board yesterday, and we've asked Joe to basically present that same material, both to the full Board today and to those of you who are there by telephone. So I'm going to turn the mike over to Joe Fitzgerald, who is with SC&A, and he's going to review for us the materials from our contractor on the Rocky Flats plant and the issues that were raised on the site profile review. Joe Fitzgerald.

MR. FITZGERALD: Thank you, Dr. Ziemer. This is Joe Fitzgerald. I led the Rocky Flats site profile review on behalf of SC&A, and good morning out in Colorado, good afternoon here. First off I just want to clarify one thing. Certainly we prepared the detailed matrix that you're looking at at this point. My presentation is essentially highlights of that matrix, focusing on issues that are significant from a dose reconstruction standpoint, or issues that would be perhaps challenges or

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barriers to doing dose reconstruction. again, we certainly did touch on, in the working group, all of the issues you're looking I believe there's 21 findings that were at. cited in the matrix. NIOSH in fact will mention it later. They have prepared a initial preliminary response to those specific issues and so certainly there's been a exchange on each and every item there, although we haven't had a chance to have a interchange discussion in a working group atmosphere. So this is, again, going to highlight what we think are the more significant issues that would be particularly important at this point in the process.

I don't believe you have a copy of my presentation, so I'm going to also cover these -- these slides I'm presenting here on the screen and more or less repeat them for your sake, as well.

The Rocky Flats profile was conducted in the fall of last year, beginning actually in early August and -- with a report being submitted after classification review on December 8th.

The Rocky Flats matrix, which you do have, is

actually the prototype issue resolution matrix that we're using for the site profiles, and it basically highlights the findings that we're looking to have a further interchange with NIOSH on and -- and looking to resolve the questions, both technical and factual, as well as those that pertain to dose reconstruction. And that matrix itself was submitted on December 15th, so that's a relatively recent review.

The first primary issue I want to touch on is the use of urine bioassay MDA median values, which may not be appropriate for plutonium and americium. And the issue there is very -- very much the -- given the rather primitive internal bioassay techniques in the '50s and even into the early '60s, our concern on the site profile is that the -- the use of the median MDA values for plutonium and americium, and even for uranium but to a much lesser extent, we feel were unduly low, given the -- given the variables involved. And the variables themselves include the counting time, the theoretical upper bound detector counter efficiency, and a number of other parameters

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which are cited in the report. But all of these contribute to what we think is a level of uncertainty that belies the estimation or the assumptions that are provided in the TBD at this point. And I think, again, it's a question of how much conservatism is really required to be claimant-favorable, particularly in that particular era of the 1950s. So what we would like to see certainly is -and I believe this is what we heard yesterday -- is NIOSH perhaps revisit those parameters, look at some of the issues that we felt were questionable, were ones that perhaps could be tightened up, and to come together in terms of what would be perhaps more appropriately conservative MDAs. And the implications of that particular issue,

I think, are important from the standpoint that, you know, as you look at the history of - of how doses were recorded -- internal doses were recorded at Rocky Flats, you know, certainly the workers on the other end of the phone can attest that in the earlier days yeah, that there were certainly practices where -- and policies where doses would be assigned. In

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this case certainly we understand that urinalysis (unintelligible) less than ten percent of the tolerance level, and the tolerance level were the -- the maximum allowable concentrations in urine -- activities in urine that would be permissible for workers at the site, and anything less than ten percent of those values would be recorded as background or zero. And that would present a -- certainly an issue if in fact the MDA values that you're using are lower potentially than those values. So we're looking at some of the implications of -- of perhaps assigning overly-low MDA values as being ones where you will miss dose. And we're concerned that at least the -- there's a need to look at that particular issue. Now in the report, as well as in the matrix, we offered forward an approach to how one could go about perhaps injecting more conservatism on these parameters. And I think the suggestion was taking two of the four parameters and using the more extreme values to come up with these higher MDA values. I think that was only offered as a -- one possible pathway. For the benefit of the folks on the other end of the

1 phone, certainly NIOSH identified another way 2 to go about looking at the conservatism and the 3 precision of these assumptions and coming up 4 with another analysis that would give you a MDA 5 value based on these more conservative 6 assumptions. Certainly we're ready to talk 7 about that and look at that. 8 I might stop for a moment. If there's any 9 questions -- we, again, had gone through this 10 yesterday so I think to some extent we had 11 covered a number of these issues. If there's 12 any questions on any of these issues -- I don't 13 think you have the benefit of any of these 14 presentation slides, either. 15 (No responses) 16 No questions, okay. 17 UNIDENTIFIED: (Off microphone) 18 (Unintelligible) we have the --19 MR. FITZGERALD: I'm talking to the folks on 20 the other end of the phone. 21 Okay, if I can turn to another issue that 22 you'll see in your matrix chart, we're 23 certainly looking at the TBD from the 24 standpoint of -- of how high-fired -- which is 25 the colloquial term for the lower, insoluble

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plutonium compounds were addressed. And our concern there, and I think we've gone through a fair amount of analysis in the report -- you know, again, were the relatively high MDA values that exist in the 1950s and '60s and certainly the -- in the .01 becquerel range -relatively high MDA value, and something that wasn't improved upon until into the mid-'60s. The low fraction of activity intake excreted through the urine was another limitation, and certainly the historic delay in or lack of post-incident urinalysis or fecal analysis were all situations where if in fact lower or insoluble, or the super S, plutonium was involved, certainly you would be concerned about perhaps missing a dose from that standpoint.

So, again, our interests or our concern on that particular issue is two-fold. One, the question of acute intakes of such compounds; and two, whether or not in fact in certain target organs like the GI tract this might be a rather significant contributor, one that has to be addressed and included in the -- in the analysis. And again, this is something that

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was not perhaps given as much attention as we would like to see in the site profile for Rocky Flats. And again, we're talking about the site profile -- and maybe somebody can correct me, I believe it was 2003 when it was generated and that -- certainly the version that we looked at, so it's been certainly a couple of years --I'm sorry, 2004, so it was a bit over a year since that was issued that we looked at it. Now my understanding based on yesterday's conversation -- discussion that we had, certainly -- and this is reflected in the report, as well -- that there's an OTIB, which is a Technical Information Bulletin, a guide -additional Implementation Guide that's being worked on and may be issued soon that addresses insoluble oxides. And this is certainly going to be the avenue by which we get more specific guidance and parameters from NIOSH to address this particular issue. Now I don't know whether it'll be in time for the discussions that we're going to have, but we're hopeful that that would be in fact the means by which we could tackle the question of insoluble or certainly somewhat insoluble plutonium

compounds.

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Another issue that certainly the SC&A team addressed is the inadequacies in neutron exposure -- excuse me one second, I'm trying to get this thing to work; there we go, went too far -- inadequacies in neutron exposure characterization. Now we've -- certainly look at the -- the neutron dose reconstruction program, one that's been underway for seven or eight years, as a good means to develop the correction factors to apply to the NTAmonitored workers, the ones that were in fact monitored using NTA film. The corrections that were made through that process I think does respond to the issues that revolved around what workers were in fact receiving from neutron exposures. Our concern, frankly, is how one can extend those correction factors beyond the NTA energies to not only other energies below the 700 to 800 keV threshold, but to other workers. Because again, the NTA film study, the NDRP* study focused on workers that worked in the plutonium operations, so it did not include workers in non-plutonium operations or workers that may have been exposed to specific sources

of neutrons beyond those production facilities. At any rate, certainly the concern is to make sure that we have correction factors that are broadly applied for neutron exposures across the Rocky Flats operations. Now at this point we feel that the NDRP data focuses on a key, but not a complete, part of that spectrum. For those on the other end of the phone, we're having some technical difficulties that we're trying to resolve here, so stand by.

(Pause)

Okay, I think we're all set here. Thank you. The other issue which I think is highly important to this particular process is that the -- and I know some of you are aware of this. University of Colorado has a number of job category-specific neutron exposure data which, in the analysis that we were involved with and the sampling that we did, certainly was very important -- important from the standpoint of looking at potential coworker models, the assignment of neutron exposures to perhaps workers that were not monitored. Now in our discussions with the University of Colorado and our discussions with NIOSH, it

just became clear that certainly the data is there, but wasn't yet available to NIOSH and NIOSH was going through great pains to try to gain access to it. So really I think our only finding and sense on this is that it's very important information and information that should be reflected in the analysis as soon as possible, but we're certainly appreciative of the efforts that have been underway to get that data.

Is there any questions on the other side? I -- are you still there?

MS. BOLLER: Yeah, this is Carolyn in Congressman Udall's office. Is that CU data, is that coming from Dr. Rutenberg*?

MR. FITZGERALD: Yes, that's correct, and there's -- again, we've talked to -- talked to him and I think NIOSH has been in regular contact, as well. So again, that -- that information is -- is certainly relevant, just a matter of obtaining and gaining access to it. And NIOSH may have a few more words to say about that since they've been in more direct contact.

The next --

1 MR. HILLER: (Unintelligible) --2 MR. FITZGERALD: -- issue I want --3 MR. HILLER: -- (unintelligible) --4 MR. FITZGERALD: -- to talk about is something 5 that we're very concerned about, and something that we feel needs to be unpacked more in the 6 site profile characterization. This gets to 7 8 the heart of the reliability or validity of the 9 data that we're -- that we're using in the dose 10 reconstruction process. And this is something 11 that certainly the Board and we, in our 12 reviews, look at quite closely at each site. 13 It's almost the cornerstone, whether or not the 14 data is reliable, whether or not we can 15 understand the pedigree for it and be able to 16 trace how in fact it is applied. 17 At Rocky Flats it ha -- certainly Rocky Flats 18 has a long history and certainly a complexity 19 in terms of the operations, and it also has 20 certainly accounts that we find troublesome 21 that we are still going through and trying to 22 take to ground. But these include a number of 23 concerns related to the data itself. Certainly 24 questions -- and this gets to the dosimetry --

questions on algorithms and dosimeter

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calibrations. That particular issue, the question of calibrating the dosimeters, surfaced both in GAO investigations and testimony on the Hill, as well as internal DOE oversight reviews. So it's not a new issue, but certainly is -- is a issue -- and this again cropped up in the '80s and '90s looking back historically -- which raised important questions about how in fact dosimeters were calibrated, were they in fact calibrated correctly and -- and what implications does that hold to the reliability of the data that was generated by that dosimetry. We have certainly other issues. The historic assignment of zero doses, null doses, and in some cases a category which I frankly haven't seen at other sites but certainly at Rocky it's been used is this question of "no data

available". And looking at the history -- and again, we -- we did a sampling review. It wasn't an exhaustive, research over time, but certainly did a sampling review. We certainly saw enough evidence that historically these values, these placeholders were in fact used in the record in a way which, in our mind, raised

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questions about the policies that were behind the use of these values and what practice was in place over time, and whether or not -- I guess the question -- the word I've -- come to me is legitimacy, whether in fact these were legitimate values or represented values that were used in place of -- of actual dose estimates or dose measurements. Again, from worker interviews as well as from the accounts, as well as the investigation reports that we looked at, more questions were raised than answered. Certainly it does speak to the reliability of the data and one that we would want to make sure that the site profile was complete on and if anything were to provide some substantiation as to how were these used, these terminologies, how these terms, these units used and on what basis were they assigned.

Another question is the presence of blanks. In this case we found that in some instances the record shows literally a blank instead of an actual reading, and in those cases we've established that at some point in time, probably prior to '64, that was used where

somebody had a security badge but did not have a dosimeter. So in fact it was left blank, and that was the -- kind of the understanding in terms of practice. Post-'64 everybody had a combination security badge/dosimetry, and at that point in time you would not expect to see, quote, blanks showing up in the record.

Now in our review we did see some instances of blanks still showing up in the post-'64 time frame. I think, again, we would raise those as issues that ought to be addressed and looked at and put to bed by NIOSH. It gets to, again, the question of reliability.

A number of these may have perfectly suitable explanations. I think our concern is that they represent loose ends on the reliability issue that need to be resolved as we go through and actually bring this to fruition.

Now the other issues I think are in your matrix so I'm not going to go through them all, but things like the placement of dosimeters in relationship to aprons and the question of where dosimeters were worn, these are issues that I think we've seen at other sites.

They're not new issues. I think, common to a

1 plant that has a history as long as Rocky 2 Flats, there certainly would be instances like 3 that. But again, we want to be sure about 4 those things and certainly the workers we've 5 talked to have raised these issues to our attention, as well. 6 7 Let me pause for a second. That's sort of a 8 lot of information. You don't have the benefit 9 to view graphs. Are there any questions on 10 some of those issues? 11 MR. HILLER: This is David Hiller. 12 MR. FITZGERALD: Hello? 13 MR. HILLER: When you -- when you suggest that 14 -- that you're not sure of the -- the reason 15 why you see blanks in the records or why values 16 were ascribed sort of as placeholders, how --17 what is the potential impact of those questions 18 if they're not satisfactorily answered? 19 MR. FITZGERALD: Well, clearly if we can't come 20 up with a substantiation of what the practice 21 had been -- you know, what you get down to is 22 some question of how you would assign dose. 23 And of course NIOSH has addressed this issue at 24 other sites and I'll let -- certainly they can 25 answer this, as well, but what we're trying to

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get to is a sense of what does this mean in terms of the data that's applied, and should in fact some special measure be recommended on behalf of NIOSH to address these gaps. They -you know, they represent gaps that may in fact be real doses, but for whatever reason, either a zero, a blank, perhaps no data available -you know, one of these terms were substituted for what should have been a -- an actual dose. We're not drawing that conclusion at all at this point, but we're just indicating that I think we need a better handle on that particular question. And if it turns out that, you know, we can't -- you know, from the historical standpoint -- figure out what the practice had been, then I think from a conservative standpoint NIOSH would have to address that issue as a gap in the database and -- and weigh the implications on the reliability of the overall data. Right now I think we're posing more questions than answers, but I think that is exactly right, we have to come to a understanding of how that happened. MR. DEMAIORI: Joe, this is Tony DeMaiori with the steelworkers.

MR. FITZGERALD: Hi.

MR. DEMAIORI: On the question of no current data available, historically at Rocky Flats that was a term used for unexplained dose. If your dose was way too high for the operation that you were currently working, they would zero it to no current data available. It was done routinely over the years.

MR. FITZGERALD: Well, obviously we would be concerned about that. But you know, again, I think -- we picked this up in the documentation. We certainly heard it from your coworkers, and it raises enough questions and concern that we wanted to convey that to NIOSH and certainly recommend that this is something that deserves further attention and -- and substantiation. At this point certainly it's -- it's one of the number of questions that go to the reliability of the data.

I want to go ahead and just speak to a number of issues which probably don't rise to the -- necessarily rise to the significance of -- of the other issues, but certainly have some potential -- some of them have some potential as we work through this. As you'll see in your

matrix, certainly we raise concerns that ranged from the assumed default particle size that was being used, and I think we had a useful initial dialogue on that with NIOSH. This is the five micron AMAD. And I think the -- certainly the notion is that there is a practice of applying what available data there is before going to the default particle size. But again, we were concerned that that wasn't as clear as it could be.

Another issue that we still haven't unpacked but we're going to spend some time on is this notion of -- of how one uses the americium 241 actual material assay as a means to calibrate against the -- the lung counting for plutonium. And our concern there, again, is where you have lung counting with either pure or relatively pure plutonium, but don't have the americium marker; is there a concern there that that may in fact pose some challenges in terms of the actual dose recorded. That's something I think we can clarify. Again, we haven't had a -- the extensive exchange yet on these issues with NIOSH. That process is just starting, but certainly we'll be talking about that.

Some of the other issues, the assignment of isotropic and rotational instead of anterior-posterior geometry, which is fancy talk for saying, you know, how is a person positioned to the radiation source. We feel that there are certainly work locations, type of work at Rocky Flats where the optimal exposure position for a worker and his badge may not be conservative capturing -- giving credit to as much radiation as possible. And I think what we heard yesterday, which is certainly very positive, is that NIOSH is considering going to a very conservative approach of using the AP or anterior-posterior geometry.

One issue that we looked at at all sites because with Department of Energy facilities there was a lot of movement of materials between sites, and you know, whether it was recycled uranium or shipments of -- of material from one site to another, this was standard practice across the weapons complex, and Rocky was no exception. And I think for Rocky Flats the recycled uranium issue was not as prominent. However, certainly over time there's questions of other materials being

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shipped, and I think we looked at two or three instances -- just as -- again, as illustrative examples, the U-233 uranyl solution -solutions from Oak Ridge. Certainly there were shipments going from Oak Ridge to -- to Rocky Flats for processing. Uranium 236 from reactor core recoveries from Idaho, that was going to Rocky way back when. So you know, you have a number of these things where we felt the site profile could be improved by characterizing some of these shipments that were not main process items, but nonetheless represented part of the operation and certainly a potential -you know, I look at U-233 with the U-232, certainly a potential for exposure for workers that might have handled that specifically. That is pretty much the highlights of the matrix as -- as we stand at this point in time. Again, this is the earliest part of the process of interchange with NIOSH on behalf of the Board. And as with other sites, we intend to engage NIOSH in these particular issues and certainly attempt to either converge or identify issues that have to be further addressed.

Is there any further questions from the audience in Colorado? Or anyone here, I guess, for that matter -- I'm sorry.

DR. ZIEMER: Well, we'll take questions here in a moment, Joe. But thank you very much. want to point out particularly to the delegation in Colorado that there already has been -- over the past I guess three months -two or three months -- ongoing exchanges involving our contractor, SC&A, NIOSH and a working group of the Board to address the various issues in the matrix. And particularly

MR. GRIFFON: That's not quite true.

Well --

MR. GRIFFON: The workgroup wasn't involved.

DR. ZIEMER: -- right, I'm sorry. There -there's been exchanges. I think -- I've seen some e-mail exchanges, at least, where the questions and so on, but we'll get some clarification here. There've been some early exchanges of information, but looking forward, we -- we have a process that does involve formal, face-to-face exchanges. It's a process that's been used for other site profiles and

1 one which will be used here involving the 2 contractor, NIOSH and a Board working group. 3 And we will want to extend the invitation to 4 someone representing the petitioners to 5 participate in that, as well. So we would let 6 you folks know when such meetings take place so that, if you so desire, you could have a 7 8 presence there, as well, as these issues are 9 discussed and we move toward resolving items 10 that are raised on the site profile. 11 Also I believe there -- the initial review of 12 the site profile was what we call Rev. 0, which 13 is the early site profile. I believe we're at 14 Rev. 1 now, are we not, Jim, as -- I'm asking 15 Jim Neton now, of NIOSH. 16 DR. NETON: Actually I think there's six 17 chapters to the site profile. Each has their 18 own unique --19 DR. ZIEMER: Oh, a revision, right. 20 DR. NETON: -- revision number. Some are at --21 DR. ZIEMER: Some are at Rev. 1 --22 DR. NETON: -- Revision 1, but most are still 23 are Revision 0. 24 DR. ZIEMER: -- right. So it's a bit of a mix, 25 but it's an ongoing process of updating the

site profile as new information is gained.

DR. NETON: Correct.

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DR. ZIEMER: Now let me open the floor for Board questions here, or comments.

DR. WADE: I'd only like to -- this is Lew Wade -- to make one comment, then maybe Jim has a comment to make in terms of the status of things with regard to NIOSH. And my comment is simply to remind you of the time lines we're looking at and sort of what's in front of us. We talked about this yesterday, but the Board does have a phone call meeting scheduled for March 14th, a face-to-face meeting scheduled for April 25th, 26th and 27th. My tentative plan is to hold that meeting in Denver, Colorado. I would certainly hope we'd be in a position to have the Board presented with an evaluation plan before that meeting and have the Board make a recommendation to the Secretary on the Rocky Flats site profile (sic). That said, there's work to be done and I would like the Board to -- to put in motion what it needs to put in motion to see that we can achieve those goals.

Jim, could you just update us very quickly as

to NIOSH's position now relative to this process?

DR. NETON: Yeah, I don't want to take up a lot of the Board's time. We went over in some detail yesterday NIOSH's draft responses to the 21 individual issues that were identified in what I'll call the consolidated matrix that came out in mid-December. It's available on the table. I believe the folks in the Colorado delegation and the union representatives or petitioners have -- have that document now. I do believe that SC&A has done their usual very thorough job reviewing this profile, and we commend them for that.

I just do want to point out that the consolidated matrix and all the efforts right now are -- we're going to try I think to be directed towards resolving issues that are of specific relevance to the site profile -- I mean the SEC --

DR. ZIEMER: SEC --

DR. NETON: -- petition we have in our hands at this time, and I think Joe did a good job summarizing the five key issues, along with the other lesser significant issues. I think the

first two he mentioned, the MDA issue and the super insoluble material, we've come a long way towards resolving or coming -- coming to terms with SC&A already. I look forward to working with them and addressing the other issues in the upcoming working group meetings.

MS. BOLLER: This is Carolyn with Congressman Udall's office.

DR. ZIEMER: Yes.

MS. BOLLER: I just want to make one statement. First of all, we believe it's extremely important that Tony and his folks be actively involved in this whole process, so it's good to know that that offer has been made and we should be kept in -- he should be kept in the loop.

DR. ZIEMER: Yes.

MS. BOLLER: The second piece is, I believe the entire delegation is very well aware of a lot of these issues, and we are very supportive of this petition. We all get those phone calls from those people who are sick and dying while we're going through this process. And while we don't want you to skip over something, we -- we would strongly encourage you to get this thing

1	done so that we can can talk with our folks
2	about it, who are sick, who worked at Rocky
3	Flats, and who need these benefits.
4	DR. ZIEMER: Yes, thank you. Mark Griffon has
5	a comment.
6	MR. GRIFFON: Just actually a couple of
7	questions maybe. One for Joe, I just wanted to
8	clarify this finding number two on the matrix,
9	and it talks about the high-fir the super S
10	class and the doses to the GI tract.
11	MR. FITZGERALD: Right.
12	MR. GRIFFON: I'm not sure we have time to get
13	into it here at the Board level, but it
14	references page 40, and all the all the
15	tables I see there compare Type S and Type M.
16	I don't see any super S compared, and it also
17	seems to me that the GI tract doses per sievert
18	excreted are higher, but it's not intake, so I
19	just
20	MR. FITZGERALD: Well, I think if
21	MR. GRIFFON: I want to know if
22	MR. FITZGERALD: Right.
23	MR. GRIFFON: there's a specific table that
24	addresses this issue of the super S compared
25	you know.

1 MR. FITZGERALD: Actually I think in the 2 analysis we established that the so-called 3 super S was synonymous with S, and I don't think we found a distinction that was worth 4 5 making the distinction for in the table itself, so the S is the super S --6 7 MR. GRIFFON: Okay. MR. FITZGERALD: -- in a sense. On the other 8 9 issue, I -- you have me at a disadvantage since 10 I don't have my internal dosimetrist. 11 MR. GRIFFON: Yeah, 'cause when I look at it I 12 see sievert per -- sievert per becquerel 13 excreted. 14 MR. FITZGERALD: Yeah, we'll have to get that -15 - we'll have to --16 MR. GRIFFON: Certainly the numbers are higher 17 for the lower large intestine and 18 (unintelligible) --19 MR. FITZGERALD: I'll have to clarify that 20 later. 21 MR. GRIFFON: -- and that would be expected 22 'cause it --23 MR. FITZGERALD: Right. 24 MR. GRIFFON: -- (unintelligible) slower, but -25 - but sievert per intake, if you -- if you make

1 sure you're bringing urine data back to intake 2 values --3 MR. FITZGERALD: Right. 4 MR. GRIFFON: -- I think -- I'm surprised by 5 that, so I guess we have to look into that 6 further, maybe at the -- Jim wants to 7 (unintelligible) --8 DR. ZIEMER: Jim -- Jim Neton can respond here. 9 DR. NETON: I might be able to clarify a little 10 bit. I don't think the super S issue has gone 11 away. I think we all acknowledge that there is 12 highly insoluble compounds, more than Class S, 13 at Rocky Flats. It's been demonstrated in 14 autopsy cases at Rocky and at the Transuranic 15 Registry. The issue with the GI tract -- we 16 believe the doses to the lung are adequately 17 covered by S because they're already more --18 MR. GRIFFON: (Off microphone) (Unintelligible) 19 DR. NETON: -- more than likely over 50 20 percent, we've been through that. The issue 21 with the GI tract is that if you calculate 22 intake based on a urinary excretion using S, 23 you're going to come up with some value. If it 24 really is super S, the intake retention 25 fraction you applied is inappropriate and the

1 intake could be possibly orders of magnitude 2 higher. If that's the case, even though it's 3 coming out more slowly, if you integrate the 4 dose to the GI tract over 50 years, you may end 5 up with a higher dose to the GI tract, or underestimate it. And that, in my mind, is the 6 7 only real issue remaining on the table. 8 I think we can pretty much demonstrate systemic 9 organs are adequately covered by S, and -- and 10 lung cancers are adequately going to be 11 probably compensated by S. So this GI tract 12 issue I think is the central issue here. 13 DR. ZIEMER: Okay. Thank you. 14 MR. GRIFFON: I agree with that. I agree with 15 that point, I just didn't see that in the 16 tables that I was looking at, so... 17 MR. FITZGERALD: Again, I think that -- that's 18 the reason. We didn't make a distinction in 19 the tables between super S and S, but focused 20 on S. 21 DR. ZIEMER: Okay. Thank you. Other comments? 22 MR. GRIFFON: Yeah, I -- this is --23 DR. ZIEMER: Mark, go ahead. 24 MR. GRIFFON: -- I guess this is for Jim, too, 25 I -- I'm just wondering, and since you

1 mentioned that -- that the priority issues were 2 focusing on now, I guess one thing that -- that 3 comes to my mind is, of the claimants in the 4 SEC petition class, is there any coworker model 5 I'm not sure whether Rocky relies on a coworker model and if that's included in the 6 7 site profile review or... 8 DR. NETON: That was actually one of the 9 comments that SC&A made was that they weren't -10 - they didn't see evidence of a coworker model. 11 And much like the Y-12 site profile, we 12 typically don't include the coworker models in the site profiles. It would be modeled 13 14 separately and we will have a coworker model 15 for Rocky Flats --16 MR. GRIFFON: (Off microphone) But since --17 since we are in this sort of middle ground here between a petition and a site profile, I think 18 19 (unintelligible) --20 We're well aware that we need to DR. NETON: have a coworker approach to cover those cases, 21 22 and we -- we --23 MR. GRIFFON: (Off microphone) And I guess the 24 same follow-up that we asked for Y-12, maybe we 25 want to get a sense of how -- how many members

1 of the cl-- what percentage of the class this 2 might affect, the coworker model. If it's 3 going to be --4 DR. NETON: Right. 5 MR. GRIFFON: -- used for 80 percent like with 6 Y-12 or is it -- is it a much smaller 7 percentage. 8 DR. NETON: Right, I think we understood --9 MR. GRIFFON: (Off microphone) (Unintelligible) 10 DR. NETON: -- that we need to give the Board 11 some sense as to what we -- you know, what 12 we're going to be extrapolating and what's 13 going to be based on real data. 14 MR. GRIFFON: And then the other, Jim -- the 15 other obvious follow-up from me, since you're 16 probably ready for this, is the integrity or 17 validity of the data. That's come up in a 18 finding. Has any of the -- have you made any 19 progress toward exploring ways to validate the 20 data? 21 DR. NETON: We're working towards that with 22 ORAU as we speak. I mean we -- we recognize 23 that this is going to be a recurring issue. 24 You see some of those issues reflected in the 25 SC&A review, but we also know that we need to

go back and look at the -- any electronic records we have and look at the pedigree of those and establish some comfort to the Board that they -- they are what they purport to be.

MR. GRIFFON: Part of the reason I raise these

is to -- to Lew's point of this time line, and these are certainly easy points to say but hard points to -- to (unintelligible).

DR. NETON: We understand, yeah, that is -- that we're under the gun right now.

DR. ZIEMER: Let me direct this question also to either Jim Neton or Larry Elliott. This is relative to the April meeting, and Lew has pointed out the desire to be ready to present a recommendation on the SEC to the Board. You have seen at least the preliminary drafts of what the Board thinks an SEC procedure for us should look like in terms of the issues that need to be addressed. Do you feel fairly confident at this point, if we have something that looks pretty much like the draft that you've already seen, that you'll be in a position to -- to address the issues, including sample dose calculations and so on, along the lines that were described in Dr. Melius's

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1 presentation on -- on the format and so on? 2 DR. NETON: We are behaving as if something 3 similar is going to be -- be the benchmark. 4 can't predict it will, but --5 DR. ZIEMER: Yes. DR. NETON: -- but we're working towards that 6 7 end, the pedigree of the data. 8 DR. ZIEMER: Yes. 9 DR. NETON: I think the second issue which is 10 key, that -- that -- again, many of these 11 issues are raised in SC&A's review, maybe not 12 in exactly the terms that are in the draft 13 document from the Board, but such -- things 14 such as if you have a lot of data, make sure 15 that all the other ancillary nuclides that were 16 there that could have resulted in exposure are 17 covered with some type of monitoring program 18 and some approach, and also the draft example 19 dose reconstructions for those activities. 20 -- but we're working towards that. We -- we're 21 trying to fill those holes. 22 DR. ZIEMER: Understood, and just -- at this 23 point no one has identified any show-stoppers 24 in achieving that, I guess, or gaps that have

been overlooked -- well, I guess if it's a gap,

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1 we don't know that it's... 2 DR. NETON: Yeah. I think the pedigree issue 3 or the reliability issue is --4 DR. ZIEMER: Is going to be the key thing. 5 DR. NETON: -- is going to be the key thing We -- we need to be moving forward 6 right now. 7 on that fairly rapidly, 'cause if we can't 8 address that issue, then the other ones are not 9 even relevant anymore. 10 DR. ZIEMER: Right. 11 I mean if you don't have good data DR. NETON: 12 13 DR. ZIEMER: Right. 14 DR. NETON: -- 'cause we're hanging our hat on 15 that data to develop these models. 16 DR. ZIEMER: Thank you. Dr. Melius. 17 DR. MELIUS: Yeah, just to follow up on that. 18 I would also hope that we'd be able, in our 19 discussion later this afternoon on the 20 workgroup report, as well as SC&A's procedure -21 - or presentation on SEC -- procedure for SEC 22 review, we would be able to meld that process 23 into (unintelligible) to the extent that that 24 would also facilitate the nature of SC&A's 25 review to not be just on a site profile, but

1	also on the SEC evaluation. I think that could
2	be helpful and (unintelligible) will be a
3	proposal to be able to get that underway right
4	after this meeting, so that's is what we
5	hopefully will be able to do and hopefully
6	that'll all work out for for this particular
7	petition.
8	DR. ZIEMER: Okay. Any additional comments or
9	questions?
10	MR. HILLER: This is David Hiller. Let me
11	DR. ZIEMER: Yes, David.
12	MR. HILLER: ask one other question, if I
13	may, please.
14	DR. ZIEMER: Yes.
15	MR. HILLER: In the the NIOSH response on
16	this on the matrix, issue nine, which is, as
17	I understand it, the issue that raises these
18	questions of reliability and validity of the
19	data, NIOSH indicates it's not the purview of
20	the TBD to correct any operational deficiencies
21	in a non-extant program. We certainly
22	understand that and don't hold NIOSH
23	responsible for what wasn't done at Rocky Flats
24	over the past decades. But it seems that this
25	goes directly to the issue that will before the

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advisory committee in terms of whether it is feasible or not feasible to estimate with sufficient accuracy the radiation dose that the class received. Am I understanding the conversation correctly that that is the -- the most critical issue that you're looking at?

DR. ZIEMER: Jim Neton is going to address that here, just --

DR. NETON: David, I didn't know if you were or were not on the call yesterday when we went over the resolution matrix, but I indicated in my presentation that we actually were off the mark in what the intent was for question -- or comment number nine, and will be revising that -- our response. The response that you see from NIOSH addresses an internal dosimetry issue, not the external dosimetry issues that were raised related to calibration and people not wearing badges and that sort of thing. Related to your -- it's not the purview of the TBD to correct operational deficiencies in nonextant programs, I'm not actually clear what was meant by that, either, at this point, and I will promise that I will get back and get an interpretation of that.

1 DR. ZIEMER: Well, you got an answer to -- that 2 is that the matrix shows the wrong information 3 4 DR. NETON: Yeah, (unintelligible) at this 5 point. DR. ZIEMER: -- so we will get the right 6 7 information and need to get that transmitted --We will be --8 DR. NETON: 9 DR. ZIEMER: -- to the Colorado --10 DR. NETON: -- revising the matrix --11 DR. ZIEMER: -- folks, as well. 12 **DR. NETON:** -- and getting that out. 13 DR. ZIEMER: Yeah. MR. HILLER: Okay. Well, I appreciate that. 14 15 But that still -- with regard to --16 DR. ZIEMER: The question may remain. 17 MR. HILLER: -- the issue of reliability and 18 validity, that still seems to be a critical 19 issue in terms of the accuracy of the radiation 20 dose. Am I right? 21 DR. NETON: We totally agree with that. Yes. 22 MR. HILLER: Let me just say then that -- to 23 follow up on Carolyn's earlier comment, that Senator Salazar, and I think all of the 24 25 delegation, is -- is focused on that issue.

1 And if we don't have sufficient information to 2 determine the dose to this class, then you 3 know, we need to get this petition resolved 4 immediately. 5 DR. ZIEMER: Yes, thank you. 6 DR. NETON: Understood. Uh-huh. 7 DR. ZIEMER: 8 MR. HILLER: Thank you. 9 DR. ZIEMER: Any further comments? 10 DR. WADE: If no one else does, I have a 11 couple. 12 DR. ZIEMER: Yes, Lew, uh-huh. 13 DR. WADE: One is a comment, one is a question 14 to the Board. I mean life with this Board is 15 about tensions, and there's always tensions, 16 and we're going to face another tension. 17 mean NIOSH, as an organization, needs to decide 18 when it's prepared to bring an evaluation 19 report to the Board. NIOSH has many things to 20 consider there. They will be considering the Board's desire for information of a certain 21 22 type. It could well be that NIOSH will feel 23 compelled to bring an evaluation report to the 24 Board before it can meet all of those tests.

And again, clearly we would rather not do that,

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1 but -- but that is a tension that we'll face, 2 and I just want it to be on the record as to 3 that tension. 4 The other issue that I have is, when the 5 working group meets -- and I assume there will 6 be a working group -- I would like the Board's 7 guidance as to whether you want those working 8 group meetings to be open to the public or not. 9 I know it's a small issue, but it's one that's 10 important for us in terms of doing our planning 11 and our noticing. So I assume that you will 12 have a working group address Rocky Flats 13 issues, and I'd like to hear a little bit about 14 that, and then I would like some guidance as to 15 whether you want those issues open -- those 16 meetings opened. If they're not opened, we 17 certainly will have the petitioners present and 18 represented, but the Board has advised 19 different ways for different meetings, so I'd 20 be curious as to your guidance. 21 DR. ZIEMER: Jim, you have a --22 DR. MELIUS: I just want to --23 DR. ZIEMER: -- comment? Jim Melius. 24 DR. MELIUS: -- respond to Lew's comments. 25 understand what you're saying, Lew, but at the

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same time I think, based on -- to some extent on our past experience and so forth, it is not helpful to the Board, to the credibility of the overall SEC process and the overall program for the premature attempt to review an SEC evaluation, SEC petition. I think it actually sets us back more than it moves -- thing ahead. It certainly tends to waste everyone's time involved, and I think if anything increases tension than decreases it. So I would hope it would be done very carefully. If there's a legitimate disagreement on -- on whether or not an issue's been satisfactorily resolved or is at the point of satisfactory resolution, that's For sort of a deliberate saying one thing. well, we don't care what you say, we're going to present it anyway, only to have us then say well, no, we want this or, you know, that's not appropriate, then -- then I don't think that's -- that's helpful and it just shifts -- it just doesn't help.

DR. WADE: I understand. I think we'll be -we'll talk about that issue a bit tomorrow when
we talk about this draft rule, the rule that's
out there and your comment on the rule. But

1	there are there are considerations that have
2	to be taken into account.
3	DR. ZIEMER: Board members, do you do you
4	have any preferences and we have we have
5	this site profile and SEC, we have others
6	coming down the line. Certainly in all cases
7	the information will be made publicly
8	available, as a minimum, and we will invite
9	representatives of the petitioners. Do you
10	want the meetings open beyond that?
11	UNIDENTIFIED: Joe, is it possible for us to
12	get a copy of your presentation?
13	DR. ZIEMER: Yes, we will we can can we
14	get that FAXed out there?
15	MR. FITZGERALD: We have electronic as well as
16	hard copy, so either way, we can
17	(unintelligible).
18	DR. ZIEMER: We'll try to
19	UNIDENTIFIED: (Unintelligible)
20	DR. ZIEMER: get that out as quickly
21	UNIDENTIFIED: (Unintelligible)
22	DR. ZIEMER: as we can. We'll need to,
23	maybe off-line, find out where to send it.
24	DR. WADE: Jason can do it.
25	DR. ZIEMER: Jason can do that. Yes, we'll get

1	that out to you as quickly as we can.
2	UNIDENTIFIED: Great. Thank you.
3	DR. ZIEMER: Board members, what you want to
4	simply declare these meetings to be open, if
5	people want to attend
6	DR. WADE: Is that Mark, I mean I Mark,
7	you've lived these. Is that a good idea?
8	MR. GRIFFON: Yeah, I mean I I think we
9	should have them open.
10	DR. WADE: Okay.
11	MR. GRIFFON: I think they'll be self-limiting.
12	You know, they I I don't think as long
13	as we're specific with our agendas, if we
14	discuss the Rocky Flats profile, the
15	petitioners and maybe a few others will join
16	the
17	DR. WADE: Okay, fine. Thank you. So I can
18	DR. ZIEMER: Can I take it by consent that that
19	represents the Board's view? Any objections?
20	(No responses)
21	Appear to be none, so we'll assume that all
22	these working group meetings will be open. We
23	will keep a record are we talking about
24	verbatim transcripts on these? Yes.
25	DR. WADE: We do that. Yes, we do.

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DR. ZIEMER: So there will be a complete and open record on all the proceedings.

Now Board members, the other issue to keep in mind as we kind of proliferate on reviewing site profiles, so far we've had one working group dealing with all of them. I know most of you would like to have that same working group do them all, and -- and the new guys are coming on board, but do you -- do you want to have at least one additional work group to sort of divvy the work up, or -- how does the exist-let me ask the existing workgroup how they feel about multiple -- I know that if you go to Cincinnati for one of these meetings, from an efficiency point of view, you may as well do two as one, perhaps. But nonetheless, it throws a burden on a few of the members to handle the bulk of the workload. Mark, you -why don't you respond 'cause you've -- you've -

MR. GRIFFON: Well, I -- we've started into
Rocky, so I sort of have a -- you know -DR. ZIEMER: You don't want to give that up.
MR. GRIFFON: -- (unintelligible) gotten into a
little bit.

1 DR. ZIEMER: Right, right. 2 MR. GRIFFON: But I mean the other question I 3 guess would be at some -- you know, at what 4 point is this a subcommittee where -- it seems 5 like it's a standing -- although we have different tasks. I guess we change tasks --6 DR. WADE: Right. 7 8 MR. GRIFFON: -- so... 9 DR. WADE: Yeah, I think we can continue at 10 least through Rocky to do this as a workgroup. 11 MR. GRIFFON: Yeah. I mean I (unintelligible) 12 13 DR. ZIEMER: Well, if there -- if --14 MR. GRIFFON: -- (unintelligible) on these conference calls and --15 16 DR. ZIEMER: Yeah. 17 MR. GRIFFON: -- (unintelligible) matrix so far 18 with Rocky, so (unintelligible) --19 DR. ZIEMER: If there's no objection, we can 20 keep the same workgroup, at least through 21 Rocky, and that again would be Mark Griffon, 22 Robert Presley, Michael Gibson and Wanda Munn. 23 The four of you are willing and able? 24 DR. WADE: Able, no question. 25 MS. MUNN: Depends on who you ask.

1 DR. ZIEMER: Okay, that will be at least the 2 working group. We can add to that if we wish 3 when we get new members aboard, but we'll start out with that. 4 The workgroup will -- will 5 coordinate with NIOSH and SC&A, and will -- we need to bring the Denver folks -- who will that 6 7 be from the petitioners there? That will be --8 MR. DEMAIORI: It'll be Tony DeMaiori with the 9 USW. 10 DR. ZIEMER: Okay. So we'll keep you in the 11 loop as we get dates established for the--12 these meetings typically will be in Cincinnati, 13 typically one-day meetings, but we'll -- we'll 14 keep you in the loop as we move forward on 15 that. Okay? 16 Thank you very much, all those out in the 17 Colorado delegation. We appreciate your input 18 today. 19 UNIDENTIFIED: Thank you. And just to add to 20 the record, let us know when there's a 21 (unintelligible) when the working group meets, 22 that'll be helpful, too. 23 DR. ZIEMER: Yes, we will do that. 24 UNIDENTIFIED: Thank you. HANFORD SITE PROFILE - PRESENTATION COMMENTS, DISCUSSION, PLAN OF ACTION DR. PAUL ZIEMER, CHAIR

NIOSH

DR. ZIEMER: Our next item of discussion is on the Hanford site profile. I would like to point out that here with us, in person today, is Livia Lam -- I think Livia -- is Livia still here? Yes. Livia is a legislative assistant for Senator Cantwell, and we welcome Livia here today. Are there others from Washington by phone? Okay, Livia, it's all on you then. Right?

MS. LAM: (Off microphone) (Unintelligible)

DR. WADE: And in your book -- in your tab on

Hanford, there is communication from Senator

Cantwell, a response from John Howard, another

communication from Senator Cantwell, so I just

point that out to you that -- so you have some

history of -- of these interactions. Under

Hanford.

DR. ZIEMER: Under Hanford. This is the -yes, this is the letter that -- it went to both
John Howard and to me, and some information was
requested, and John Howard went ahead and
answered that, and I saw no need then for me to
answer it, although I still could follow-up if
the Board so desired. But this was -- this was

one that was attended to. Does the Board feel there's any additional correspondence necessary at this point? Recall, again, I could not respond without the Board's input, but since it was addressed to both John Howard and to me and John provided the requested information, we basically let it ride at that. And you have the correspondence there.

Well, let us proceed then with the --

DR. WADE: No, I need to do my conflict -DR. ZIEMER: Oh, yes, we need to do the
conflict of interest statement.

DR. WADE: Now this is a site profile discussion, and my notes would indicate that the only Board member conflicted is Wanda Munn on Hanford. Let me point out to you that the policy operating to is that when discussing a site profile, Board members who have a conflict may participate in the discussion at the table, but cannot make motions or vote on motions. So Ms. Munn, you are -- you are so instructed.

MS. MUNN: I understand.

DR. ZIEMER: Okay. So the presentation on Hanford for the site profile review will be given by John Mauro. John, welcome.

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DR. MAURO: Good afternoon, everyone. Ziemer, Livia, nice to see you again. I'll be giving the presentation on Hanford. In fact, I think we should set the table a little bit for Hanford because it was about a year ago -- a little less than a year, maybe 11 months ago -when work actually began. It was one of the first site profile reviews that we began, and it involved the typical steps -- you know, reviewing the document, and we had a large team of people reviewing the document, reviewing all the references. We did meet -- we did meet with many representatives, workers, experts. We did have extensive discussions with NIOSH, authors of the site profile, and the work product came out -- it's this 291-page document that was issued in June. And since June a lot has happened. A lot of the issues that we raised here, there are -- and by the way, we did -- let me move this along. Let's see, which one -- I guess I should use this right here -- it's the other one now, and I guess just press the top button or the center button? We'll see in a minute. No, not that, let's try this.

(Pause)

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Okay. As I was mentioning, on June 10th the large site profile review was issued. Then we were requested -- then we moved into this mode of preparing matrices. You know, a 291-page document is a lot to lift and digest and to use as a tool to try to track things, so the Board had requested on November 16th that we prepare an issues-tracking matrix, and that of course is the way we are managing ourselves these days. And we did, very recently, on January 16th send out a two-page matrix that summarizes the 11 issues -- major issues as -- it's -- to try to boil down a 291-page document into 11 issues, but you know, I think -- you know, it serves its purpose, as long as we, you know, don't lose sight that there's a lot of texture here. And so what I'm going to be doing is going through the matrix.

What I'd like to point out is when -- when you -- when you step back and you look at the site profile, you find out that it really boils down to major concerns. There are a lot of concerns, but our major concerns have to do with neutron doses, especially in the early

years, and we're going to talk about that quite a bit. And the other has to do with exposures to some exotic -- I call them exotic -radionuclides. Because of the very complex array of transuranics and radionuclides and experiments that were involved in this -- at this site over many, many years, having a good handle on the types of radionuclides that were at play and what the doses were to people who were involved in handling that material becomes very challenging. So we're really going to be talking about external exposure, neutron; internal exposures to plutonium, americium, curium and many other radionuclides that are -are of concern for the reasons we'll -- we'll get into.

Neutron exposures are important for a very simple reason. If you think back to the reactors in the 100 area, they basically had a neutron reflector and they had a thermal shield, a biological shield. But then they had hundreds -- many hundreds of holes, ports, in which instrumentation, fuel was entered and you look into the record, you find out that there was -- lot of maintenance had to be done on

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these units and that -- you also look into the record and you find out many of the workers had -- they determined they had some sodium 24 as a body burden, which means that -- that's -means they were exposed to some neutron exposure which resulted in neutron activation. So our main concerns sort of, as we got into the process, is that exposure to neutrons, especially in the early years, and they were not really adequately monitored, especially in the early years. In fact, a good break point is 1972; 1972 was the year when they moved into the Hanford multi-purpose dosimeter. That's a dosimeter that has five elements, uses TLDs and does a very nice job of measuring not only gamma exposure but also neutron exposure. Now we have lots of comments on that, too, but those are what I call second order comments where you could deal with -- you know, coming at adjustment factors for properly interpreting that data.

But prior to that -- or our main concern is that okay, how good a job was done in the -- in the -- in those early years when they didn't have the Hanford multi-purpose dosimeter?

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Well, in those early years what was done was take advantage of data that you had on the neutron-to-photon ratios. You can almost visualize one of these pass-through reactors or -- in the 100 area with all the ports, and whenever you have a stream or a photon field, you're going to have some neutrons coming along with them. The -- and so -- so if you can get a handle on the ratio of neutrons to photons, you've got your first hook into coming up with the neutron dose. So what -- what -- our main concern was was a -- was a good job done in coming up with the neutron-to-photon ratios. Now -- and we're going to get a little -- now we dig down to the weeds a little bit because this is where the rubber meets the road on this particular issue. Okay? We're going to go back -- between 1950 and 1961, this was before the multi-purpose dosimeters, the single-pass reactors, and I listed the reactors, there -- there's good reason to believe that the workers there experienced some neutron exposure. They wore NTA film and they did see positive readings. They wore their regular film badges where they

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got photon exposures. They said well, listen, we need to somehow get a ratio, neutron-to-photon ratio with the data we have.

Now the first thing that we notice is that with the way in which the site profile comes at the problem, it says well, we found seven workers that worked at those facilities at that time where we have both NTA film -- neutron detector film -- and -- regular film badges, and so from that you can come up with a neutron-to-photon ratio, so we've got seven workers. Well, this was the first -- now that one piece of information is the first place where our antenna goes up, says hmm, you've got all these workers working over all these years -- I don't know how many, there may be literally thousands of workers somehow involved, let's say, in working with all these reactors. We're really not quite sure what the number of workers, but in the end what we're saying is we have data for seven workers where we have neutron-tophoton ratios, where you simply look at the photon exposures of the film badge and the NTA exposure and you get a ratio of one to the other. And we have it for seven workers.

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Now what NIOSH did, and ple-- and by the way, this is the first time we're bringing all this forward. We have -- this is -- this may be day one of what we would call the review cycle, because prior to today you really haven't entered into a dialogue. So as I describe -summarize this information -- be very interested in feedback as to whether this is the proper understanding of what in fact was done in this -- by -- in the site profile or whether we -- perhaps we got it wrong, but my understanding, our understanding is for those years, for those reactors, what was done to come up with the neutron to photon ratio was they had seven workers. That's not very many workers, given the complexity and size of the site to number of workers. So that -- that's -- I guess we'd say issue number one, is that good enough. Second thing is they said well, we've got five

Second thing is they said well, we've got five different -- we have -- we have these seven workers, but there are a lot of different ways you could come up with this neutron-to-photon ratio, and -- and as we understand it, NIOSH used five different methods. On the left-hand

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side is the most conservative method and on the right-hand side is what we'll call the most realistic method. So you could -- so the neutron-to-photon ratio, based on these seven workers, the average, went from a high of .431 -- which means if you have one rad from -- from gamma, you've got .431 from neutron -- and down to .09. Well, what -- what I -- our understanding is, in light of this and the uncertainty, the decision was made, let's go with this distribution. So in other words, if you're going to reconstruct a dose to a worker and you have some film badge data and you say I want to try to figure out what his neutron exposure is, you multiply his film badge readings with its uncertainty by this distribution, and now you've got a distribution on the neutron dose for those -- for those workers.

Now -- so that's our starting point. It's almost like the rock we're standing on, at least for those reactors at that time period.

Okay. Now -- but then they said well, there's one more thing we have to do. In addition to coming up with this distribution, we recognize

that NTA film is really not a very good detector, especially for neutrons that might be coming from a reactor where the neutrons are attenuated and they may actually be attenuated down to an energy level that's below the threshold that can be seen by NTA film. So what happened was the -- NIOSH decided that well, we believe that film badge -- NTA film that they had data for really only captured 28 percent of the exposure, so they multiplied that distribution by 3.57. Okay? And now they have an adjustment factor that says now we're going to -- to account for the fact that the -- the neu-- the NTA film itself has these deficiencies.

Now when we look at that, we ask ourselves well, here's where our second -- the antenna go up again. We say to ourselves well, is that -- is that good enough. Is there any reason to believe -- you know, they did not have -- as best -- to our knowledge, the actual energy distribution of the neutron flux coming out from these reactors for different locations, different times. So in order to come up with that adjustment factor, you have to have some

knowledge of what the energy distribution is of the neutron flux that you're dealing with.

Now if it turns out that the vast majority of the neutrons leaving the reactor were below .7

MeV, well, you're really not -- for all intents and purposes, you're not going to see anything.

So -- and if you don't -- so that becomes an important factor. So when we enter into our dialogue and -- with Jim, you know, we're -- we're going to be talking about whether or not that .28 -- or the 2.57 multiplier is a good number or not.

Now from previous meetings we heard a lot about well, now -- well, the reality is you can see .3 MeV. I -- I would argue that that's a little misleading. I'll explain why.

If the proton in the gel that -- in the film -- in the NTA film is moving with .3 MeV, ripping its way through, it'll create enough tracks that you will be able to count a track. But for that proton in the film badge to experience a 3 MeV, it's got to be hit with a billiard board -- ball head-on collision, right square on where a 3 -- .3 MeV neutron crashes like a ball -- like a billiard ball, crashes directly

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into a proton and delivers that full -- its energy to it and it takes off. If it's a -- if it's a glancing blow, and it turns out only a very, very small percentage of the neut-- of the neutrons that interact with protons are these direct hits, so yes, someone could argue in theory you could see something at .3 MeV if you have a direct, head-on collision, but most of the collisions aren't. So from a practical standpoint, and here's a place where we'll get into a dialogue, our position is well, for all intents and purposes, if you're below .7 MeV you're going to start missing an awful lot. So issue -- you know, the second issue within the -- this -- what I call the neutron-tophoton ratio issue for this time period is can we really -- is -- is it good enough to use the measurements from the seven workers. is the .28 -- 28 percent adjustment factor for NTA film, is that good enough, does it really give -- is -- is it doing justice to the situation really. We'll move on. Let's move on to the N reactor 'cause this is a little later generation. See, those were the earlier reac-- pass -- one-pass reactors.

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we have a closed-loop reactor. Now what happens here is -- we have the same problem. We don't -- we have to come up with what the neutron-to-photon ratio is, but now we're dealing with the N reactor. When -- when all -- when all is said and done, what we found out is well, they basically used the same data, the -- the same information, except they threw one more factor in. They said well, we know we -we believe -- we believe that for the N reactor -- we have all this experience from these single-pass reactors, and we probably need to fix some of the -- the problems we've seen in these other reactors where there was -- there were lots of problems regarding neutron and photon exposures, so there's literature that goes in there -- goes into a description of shielding that's going to be added to this generation of reactor, or to the N reactor. And on the basis of that analysis, a reduction factor I believe of -- a seven-fold reduction factor was applied to the -- to the distribution. Remember the distribution we had before? Well, what they did is said well, whatever that neutron to photon ratio was, the

one we had before, we're going to reduce it by a factor of seven because we're going to take credit for the additional shielding that was put in.

But one of the things that we found out, and this is a question that we need to discuss, is we don't know if that shielding was ever really installed. They talk about it. There are calculations about it would be nice to have this shielding to correct this problem. But right now one of our questions is was that shielding ever installed, and did it in fact achieve a seven-fold reduction in the neutron-to-photon ratio. Okay? So that's -- that's an issue.

Let's go on. Oh, now we're going to -- we're still talking about neutron-to-photon ratios, but now we're going to talk about the plutonium facilities, plutonium finishing facilities in the 200 area. What they -- they took a different tact (sic) here -- okay? -- in the TBD, as we understand it. Post-1972 they had the Hanford multi-purpose dosimeters, which means that they got some really good measurements, neutrons and photons, so you

could come up with a good neutron-to-photon ratio. And they did. They said for post-'72 data we'll -- we'll take that data and they came up with well, we know what the geometric mean -- the geometric standard deviation for the distribution of the neutron-to-photon ratios are, so now we have a real good handle -- if we know what the photon exposure is, we could use this distribution to get a handle on the neutron exposure. And the -- and the multi-purpose dosimeter captures the full range of energy, so it's a good -- but there are -- I don't want to discount -- we do have some concerns about that, also, but those are what I call second order concerns.

Now, the fir-- you'll see that -- our comments on this, we have basically two comments. One is we notice that when they did that work -- that is, come up with those ratios from the HMPDs -- they went with a dataset where the measured values were above 20 millirem, but we noticed that the minimum detectable limit was 50 millirem, so I'm not quite sure how that, you know, plays out. That is, if -- if they selected set -- datasets that -- where they

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said -- where -- that these are the -- the HMPDs that we're going to use and they picked -- they selected them based on a cut-off of 20 millirem as being a low limit of detection, but the actual low limit of detection for neutrons was 50, I'm not qui-- I'm not quite sure and we're not quite sure what implications that has with respect to that distribution. But that -that, I would say, is a relatively minor issue. The more important issue that we -- we are concerned with is they're taking that ratio now that they obtained from post-1972 data and they're going to apply that ratio to pre-'72. Okay? That sounds reasonable, everything else being the same that's a reasonable thing to do. But we have reason to believe that everything else wasn't the same. That is, in the earlier years there -- there's literature that says that -- that there was a lot more hands-on operations. In addition, there was a lot of additional shielding installed. So the question becomes the distribution of values, the neutron-to-photon ratios that we see post-'72, is it appropriately applied to pre-'72 without any adjustments to take into

1 consideration some design and operational changes that occurred.

Okay, that covers the neutron issues, and I would imagine the -- we will be having meetings, and we're going to be talking about it. But it's really clean. I mean the issues -- you know, the areas where we have our concerns, there's -- are something you can sink your teeth into so I'm expecting that when we do engage in this process what we'll -- we'll --

DR. ZIEMER: John, before you leave the neutrons, I'm a little surprised -- I don't know if you or Jim can speak to this, but if you'd asked me just out of the blue, I would have guessed that the Hanford folks had spectral data in workplace for neutrons. Did they not have spectral data? 'Cause I would have thought the practice was that they would take the spectral data and then have a calibration factor against either the NTA film or whatever they're using to -- to go from whatever the film detected to dose. Is there no evidence of spectral data in the workplace for these various reactors?

1 DR. BEHLING: I think that's explained in our 2 TBD review, and the 28 percent or the 3.52 3 correction factor actually used a tissue 4 equivalent proportional counter as a reference 5 value to NTA film. In other words, take the 6 tissue equivalent proportional counter in a 7 given facility, measure its dose or integrated 8 dose over a period of time and compare that to 9 NTA film, and that's where the 28 percent 10 correction factor comes into play. At least 11 that was the way I interpreted it when I read 12 the TBD and -- and therefore the issue of --13 DR. ZIEMER: That would have been specific for 14 a particular facility then. 15 DR. BEHLING: Well, it is specific for the 16 eight single-pass reactors that were identified 17 in one of the slides that John showed where you 18 had the seven workers and there were the five 19 methods --20 DR. ZIEMER: Okay, so that's where that came 21 from. 22 DR. BEHLING: Yes, and it's --23 DR. ZIEMER: Okay. 24 DR. BEHLING: -- it's really not based on

spectral analysis, but comparing one NTA film

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1	against the tissue equivalent proportional
2	counter.
3	DR. ZIEMER: Okay. Sort of indirectly that is
4	a spectral
5	DR. BEHLING: Yes.
6	DR. ZIEMER: analysis. Jim, do you add
7	to that or
8	DR. NETON: No, I think Hans Hans has
9	addressed the issue.
10	DR. BEHLING: Good.
11	DR. NETON: I just there's a couple of
12	things with neutrons, though, I'd like to point
13	out. One I think this issue of sodium 24, that
14	doesn't necessarily mean it was an activation
15	product in the body. There was actually sodium
16	24 in the drinking water.
17	DR. MAURO: In the drinking water?
18	DR. NETON: Yeah, I think so. That that's
19	not very good evidence
20	DR. MAURO: Okay.
21	DR. NETON: at all.
22	DR. ZIEMER: Could be, yeah.
23	DR. NETON: And the second issue I think is
24	when you were talking about these detection
25	limits, you've got to remember you're doing

1	you're generating distributions. And as we've
2	talked about many times, when you're generating
3	a lognormal distribution, the detection limit
4	really is not relevant as long as you're rank
5	ordering doing cumulative probability plots,
6	you can still pick off the 50th percentile in
7	the geometric standard deviation. So whether
8	it was 20 or 40 is not really a central issue,
9	I don't think.
10	DR. ZIEMER: Yeah.
11	DR. NETON: If you're doing geometric
12	DR. ZIEMER: Yeah.
13	DR. NETON: if you're doing lognormal
14	distributions, the detection limit is not
15	really relevant.
16	DR. ZIEMER: Well, these are items of course
17	DR. NETON: For generating lognormal
18	distributions.
19	DR. ZIEMER: that'll come out in the
20	DR. NETON: We can talk about these
21	DR. ZIEMER: in the exchange
22	DR. NETON: technical issues, I just wanted
23	to point out
24	DR. ZIEMER: it just I was just struck,
25	as you

DR. NETON: Okay.

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DR. ZIEMER: -- went on there that it seemed to imply that they had no spectral information and it astounded me because I know who some of those HPs were there in the early days and I was just certain that it existed somewhere.

DR. MAURO: Okay.

DR. ZIEMER: Okay.

DR. BEHLING: Just another comment that I guess could have been added in John's presentation and that is the issue of the seven workers and the five methods were basically derived on the basis of NTA film. And I think there's a certain paradox to that because in one instance we say that NTA film was not reliable to monitor people, but at the same time we're saying it's good enough to measure neutron-tophoton ratios. So one should also keep that in mind when you look at the pedigree or the reliability of establishing these five methods and deriving from that the neutron-to-photon ratio.

DR. ZIEMER: Thank you. Thank you, John. ahead.

DR. MAURO: Okay. We're going to move on to

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internal dose, and this sort of presents an overview of the issues related to internal dose, but we're going to get into a little bit more detail in the next few slides.

I guess the -- from a priority point of view, recycled uranium is raising its head again, as it always does, as being an issue, the ade-- or the ability to fully appreciate what the internal exposures may have been to recycled We're talking about thousands of uranium. workers that processed and handled RU, and there are default values used in the site profile, and I guess the bottom line is can we hang our hat on those ratios -- that is -- and I think some work needs to be done and a dialogue engaged on whether or not the default values -- not -- capture the full distribution of recycled uranium and the trace levels of plutonium and americium and neptunium in a way that accounts for the uncertainties.

Bear in mind that it really is not until 1988 when you have alpha spectrometry on neu-- so prior to 1988 when we -- get into that right now, you've got yourself a situation where you really don't have great data, great bioassay

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data upon which to get a full appreciation of what the intakes were of these various radionuclides. And so the neutron -- so the recycled uranium is one issue.

But then there are all of these what I quess we call special campaigns. As we read through the supporting literature we find out that there was an awful lot of experimental work going on. One of course that was mentioned over and over again is the use of uranium-233 as fissile material, and its associated uranium-232. also have come across that apparently there were -- experimental work being done where there was large quantities of cobalt-60, carbon-14, yttrium, polonium-210. So what I am getting at is that there's a richness of mix of radionuclides that have occurred at different places at different times that right now we're un-- we're not completely convinced that it's been aired and the uncertainties aired adequately in the TBD. I think this is a subject that we need to engage and discuss how well are we able to track and reconstruct doses to these categories of workers. Can we identify these categories of workers that may

have experienced these exposures, how extensive it was and can we get a handle on it as to -- and in many respects it's like trying to trace down some of the issues we had when we were talking about Mallinckrodt. There were other radionuclides. There were other -- that I think -- that is not fully captured in the TBD and we -- and we'd like to engage NIOSH in a conversation regarding that.

Now, given that you have this uncertainty regarding what in fact was the internal exposures, we -- you take a close look at what was actually done. What is -- what are the instructions given to -- to the dose reconstructors and how -- how well do those instructions hold up. Now when you don't have any bioassay data available -- that's the question. I mean if you have bioassay data available and you can trust it and it's complete, great. But when you don't have any bioassay data available, that's when you have to start to use surrogate approaches. Now in the TBD they basically cover three different time periods. Say well, from 1943 to 1946, before there was any bioassay data, what

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we're going to assume, there's reason to believe that a worker may have experienced some internal exposure -- entered an area with airborne activity, what we're going to do -and on face value this sounds like a very reasonable and claimant-favorable approach -for that time period -- we do know that if the levels were above a given quantity in terms of dpm per cubic meter, respiratory protection was used. So you know what -- what's going to be -- what's -- their -- decided to do is say what we'll assume is that anyone that had a potential to be exposed to airborne activity in that time frame, '43 to '46, we will assume that he experienced exposures whereby the distribution of the airborne radioactivity had a triangular distribution that went from zero to a mode which -- which was the level that was required for respiratory protection to twice that, and that he went in there without respiratory protection. So that was the sort of the fall-back position. When we don't have any information, we'll just make that assumption.

Then from 1947 to 1952 they changed the

approach a little bit and they said what we're going to do is we'll simply assume that when a person does enter an area that -- where he may have received some internal exposure, we'll assume he's at ten percent of the level that you're required to wear respiratory protection. And finally, from '53 to '88 -- '88 is an important date because that's when they went to alpha spectrometry on the urinalysis so you got a good handle, but you had to fill it in from 1953 to '88. What they simply assumed was that if a person enters such an area they were at the -- ten percent of the allowable limiting concentration for either strontium-90 or plutonium-239. So this becomes the one-sizefits-all.

Now -- and I have to say, as a health physicist looking at that, say you know, that's a pretty conservative set of assumptions to -- you know, to come at the problem that way, saying that everyone, you know, is -- that had any potential to be exposed to airborne activity. But then -- then you have to think about it a little bit. At that time, when they were taking those air samples, were they in a

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position to make a good -- a reliable judgment of -- that in fact the levels were at some fraction of the allowable limit, especially considering the fact that a full appreciation of this full array of radionuclides that we're dealing with was not disclosed. So were the -here's the question that we need to talk to NIOSH about. The data that was available to the health physicists in 1943 to 1946 time frame, did that put them in a position that they had a good control over the situation where they knew that in fact -- whether or not there was a problem or not, especially when we talked about other issues, the breathing zone versus general air samples. As you recall, there could be a ten-fold difference there. So there -- there are issues related to the adequacy of the knowledge at the time of the -regarding the air samples to be able to make that judgment whether you needed -- whether or not we had a problem in an area or not. Now what confounds this further, and this is item number two under SC&A comments, is confirmation of the adequacy of these assumptions is very limited because there was

abso-- there was no bioassay program from '43
to '46, according to the TBD and the supporting
literature. Bioassay started for plutonium in
'46, for uranium and fission products in '47,
and by late 1960 there were -- they were -they were doing some more advanced bioassay
studies, but it wasn't until '83 -- not '84,
'83 -- that we had a alpha spec. So the
question becomes does this one size fit all for
a way of filling in this incredible gap. For
internal exposure, is it adequate. And that's
a question that we'd like to discuss with -with NIOSH.
This is the last slide. There are a number of

This is the last slide. There are a number of other issues that -- that we call the lesser issues, but nevertheless issues that we don't want to lose track of. One is something that we talk about quite a bit that might right now have been solved. We found when we read the TBD that there was some ambiguity on the instructions on how do you interpret the data that you have, the bioassay -- not bioassay, the film badge data or the TLD data. How do you assign uncertainties. And this goes back to discussions that Kathy Behling and Hans had

regarding confusion, because in many cases we found -- I use the word impenetrable, the information is so complex that we fou-- we are finding that the folks that are actually doing the dose reconstructions are very often confused. But now that we have the workbooks, we think a lot -- that might -- the problem might have gone away.

There are these oth— there — there are issues that we — we notice that — not — there was not a great deal of attention given to extremity dose, skin dose, gonad, breast, to the — you know, the beta emitters and the weak photons in the TBD. We'd like to engage a little bit more and learn a little bit more about how you deal with that.

There was also -- we noticed that for environmental exposure -- this was an interesting observation, but it's not -- I don't think it has a big effect on doses. And you -- visualize you've got these emissions coming from the various release points in the plant and you've got workers -- unmonitored workers outdoors. And you want to try to put an upper bound on the exposures that those

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workers may have experienced. Well, what they use is they took the source term information that I believe came out of some of the dose reconstruction work and they applied atmospheric dispersion factors using a computer program called RATCHET, which is a very good program and it does what's called puff invection modeling. But we noticed that they didn't use the puff invection feature. just used as average annual (unintelligible). So we -- we think that there's room for improvement on how to do that kind of analysis. And finally -- and I don't want to give it short rift (sic), but I realize I'm taking up a lot of time, we felt that there's more analysis -- more discussion is needed of exposures associated with the tank farms. discussion is needed with regard to exposures during D and D operations, which was quite extensive. And finally, there were an awful lot of incidents that occurred, and there really is very little -- there's guidance in -and this is a recurring theme. You know, right now we're depending on the CATI interviews to lead us toward whether or not a given worker

may have experienced an exposure from an incident. It seems that a little bit -- more needs to be said for guidance to the dose reconstructor regarding incidents and making sure that -- that they don't miss an exposure from an incident.

And I think that concludes my presentation.

DR. ZIEMER: Okay. Thank you, John. Board members, do you have questions for John at this time? Wanda.

MS. MUNN: Just a couple of comments rather than questions. Thank you, John, for a very good presentation on an extremely weighty site profile. It is a hard document to get through, and I'm pleased to see that you've refined the issues to things that I think probably you and NIOSH can ultimately work down to only one or two major issues for addressing. It strikes me that the work that both NIOSH and you are doing in this regard is based on the work of the men and women who established health physics programs at this site during those early years, and were -- I think we're fortunate at the Hanford site to have had a continuum of record-keeping that made it possible for a depth of

1	information that isn't perhaps available on all
2	other sites to be accessible to you.
3	I don't want personally to comment on any of
4	these issues until after there has been more of
5	a dialogue between NIOSH and SC&A. This is the
6	first time I have seen this this
7	information. I don't even have the hard copy
8	yet. I assume we will get hard copies. Right?
9	DR. MAURO: Of the matrix or of the report?
10	MS. MUNN: No, we have the matrix.
11	DR. MAURO: Oh, the the I put 20 copies
12	out. I should have put out more. They're
13	they're all they're all gone.
14	MS. MUNN: Okay.
15	DR. MAURO: I have we can make up more.
16	MS. MUNN: Fine.
17	DR. ZIEMER: I don't think we got copies, did
18	we?
19	MS. MUNN: Huh-uh. Thank you.
20	DR. ZIEMER: Okay, we need to get copies.
21	Okay, thank you.
22	Let's see, Jim, do you have any comments at
23	this time you want to make on
24	DR. NETON: No, I'd like to thank John for a
25	very very concise presentation. You know,

we -- I'm glad that we've been able to condense a 290-page review down into something that's a little more digestible. We look forward to working with -- with SC&A on this. We have just seen these issues raised today. I mean we knew they were in there, but they were in the 291 pages and I'm glad we're focusing on these and, through the workgroup process, I assume that we'll get together and come to some resolution.

timetable-wise, what we're -- where we need to be on this in terms of target times?

DR. WADE: Well, I think the driver is that, you know, we've had -- this is one of the first reviews we've commissioned. I think it's -- it's been in the process of getting to the Board, so I think there's an urgency that comes from that. I know of no other urgency, but I think it's important that we resolve these issues as quickly as possible.

DR. ZIEMER: Lew, can you inform us on --

DR. ZIEMER: I'm trying to get a feel for the extent to which -- in the context of the Pacific Proving Grounds SEC and the Rocky Flats SEC, that we'll be in a position to -- to act

1 on this in the next couple of months. 2 UNIDENTIFIED: (Off microphone) 3 (Unintelligible) 4 UNIDENTIFIED: (Off microphone) 5 (Unintelligible) DR. ZIEMER: And Y-12 of course, yes. 6 7 DR. WADE: I think we need to -- from my 8 perspective, I think we need to get NIOSH and 9 SC&A together, the sooner the better, to try 10 and deal with these issues. I assume we're 11 going to have a Board member or two present --12 DR. ZIEMER: Yes. 13 DR. WADE: -- when that happens. I think that 14 would probably grow into a working group, so 15 you know, I think there are things we can do. 16 DR. ZIEMER: Okay. Mark. 17 MR. GRIFFON: I guess that would lead into my 18 next que-- I mean this is not really a Hanford 19 question, but it's a -- just a question about 20 the upcoming workload and -- for -- for NIOSH, 21 for SC&A, as well as the workgroup -- Board. 22 I'm thinking about Rocky Flats, Y-12 and PPG. 23 DR. ZIEMER: That's what I just mentioned. 24 MR. GRIFFON: And you know, a couple things 25 that I -- I wanted to ask about -- we're in the

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middle of the Y-12 site profile review and we were talking on the side here about this CD with the 6,000 pages, and I guess the question that arises in our mind for -- with the limited time frame coming up or -- or at least we're trying to come to some conclusions as quick as we can on this, we -- we were just having a discussion of how best to proceed with reviewing the material on that -- on that CD. I guess the concern is that it may take NIOSH several weeks to -- to digest this, get it in a database format and then turn it over to SC&A. So I guess I just wanted to maybe discuss that as a -- it could be a separate task entirely that -- that we might want SC&A to, you know, sort of be gleaning this data in parallel with -- with NIOSH instead of in series, you know. So it's just something that came up, and the other thing that we haven't really yet -- we mentioned that NIOSH should get together with DTRA and -- and discuss the PPG issue. We did ask that Board members be involved. I wonder if we need to have an SC&A presence there, too, or -- or are we going to task SC&A with assisting us in that review. And that doesn't

1 seem like it would be a lengthy item, but it 2 could be a time-sensitive item, certainly, and 3 it may involve -- I hope not, but it may 4 involve our classi-- you know, classifying 5 groupings or meetings or whatever, so --6 DR. ZIEMER: Okay. 7 MR. GRIFFON: -- just -- just a few --8 DR. ZIEMER: Okay, well --9 MR. GRIFFON: -- things to think about. 10 DR. ZIEMER: -- for now, let's cogitate on 11 that. We can revisit it tomorrow if you want 12 formal action. But the point is that there are 13 a number of issues that are quite pressing. 14 may be three or four weeks before NIOSH has a 15 chance even to react to this in a formal way. 16 I'm just trying to get a feel for whether or 17 not we need to formalize any process for the Board at this point until that occurs, and then 18 19 perhaps at our phone meeting see where they are 20 on that and determine whether we need to set up 21 a workgroup at that point. 22 DR. WADE: Well, I gue--23 MR. GRIFFON: Yeah, I think Lew said it best, 24 that -- that -- let NIOSH and SC&A talk, and 25 then it may -- it eventually will evolve into a

1 workgroup function I think. Right? 2 DR. WADE: I would like to think that it'd be 3 an interaction. Now whether you want that 4 interaction to take place with a Board member 5 present on the phone call, that's your choice. But I think there's an opportunity to start to 6 7 make some ground on the Hanford issues and I'd 8 like to see us take that opportunity. 9 DR. ZIEMER: I think based on past experience, 10 the Board's preference is to at least have 11 someone present at these, and we can so assign, 12 if we know when -- when the exchanges will take 13 place. And then, again, as has been the 14 process to copy the Board on e-mail exchanges 15 that occur on issues of this type. Is that 16 suitable? 17 DR. WADE: It's fine for me. 18 DR. ZIEMER: Yeah. 19 DR. WADE: Would it be best to identify a Board 20 member now or do we want to do that -- think 21 about that this evening and talk about it 22 tomorrow? 23 DR. ZIEMER: Let's talk about it tomorrow. 24 DR. WADE: We have some new members, too, so --25 DR. ZIEMER: Right, an opportunity to chat with

different ones. Okay.

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DR. NETON: I'd like to suggest -- in the past what we've done is to provide NIOSH a chance to react to these -- you know, these more refined comments and -- and maybe just engage in a technical discussion where need be with SC&A --I mean a Board member would certainly be invited to attend, but not necessarily present because these would be at a sort of -- fairly detailed technical level. As John was talking today I could see we're going to get down into the weeds on these issues and -- and SC&A in the past has been pretty good about preparing minutes of those discussions that would be available to the Board.

DR. ZIEMER: Right.

DR. NETON: And then at least we could get some rough draft responses from NIOSH fleshed out and available for further discussion.

DR. ZIEMER: Right. That certainly has worked well in the past, and no reason not to continue that -- that mode of operation. It doesn't preclude these early exchanges just between the two of you.

DR. WADE: And so then we might have some

interaction between NIOSH and SC&A on these
issues to try and -- a clarifying call, the
first step, and we would -- we would ask SC&A
then to see that there was a transcript of
those interactions and a summary provided to
the Board. I think that gives us --

DR. ZIEMER: Right.

DR. WADE: -- a step we can take.

DR. ZIEMER: Good. Okay. We're going to go ahead and take a break at this time. We have a 15-minute break scheduled and then we will resume at that point. Thank you.

(Whereupon, a recess was taken from 3:20 p.m.

to 3:40 p.m.)

REPORT FROM SC&A ON SEC TASK DR. ARJUN MAKHIJANI, SC&A

DR. ZIEMER: We are ready to proceed in the next item on our agenda. Now if -- if you would look at your agenda a moment, you'll notice that there is an item called "Procedures for Board Evaluation of SEC Petitions", and then you'll see after that "Report from SCA on an SEC Task". What the Board would like to do is reverse the order of those two. The Board would like to hear the SE-- or hear the SC&A report before it holds its own discussion on

the SEC procedure so that we can use the SCA -don't you like these acronyms -- so we can use
the SCA information to inform our discussion of
our own SEC procedures.

So what that amounts to is the following, that
-- the Board has recently developed what we
call Task Five, which is a newer task for our
contractor which allows our contractor to get
involved in assisting us in various aspects of
Special Exposure Cohort reviews.

Under that task there were two initial subtasks that were assigned to the contractor. The first of those, subtask one, was to ask the contractor to review the procedures used by NIOSH and by NIOSH's contractor, ORAU, on the SEC evaluation process. So we have that report that we've received from SEC (sic). And Board members, that -- the report itself -- dated November 23rd, and there is a copy in your notebook.

And then subtask two where we asked the contractor to give us its thoughts on how the Board itself should proceed in handling SEC petitions, what our procedures should be. And of course parallel to that you recognize that

the Board has had its own working group that has developed some SEC criteria by which we will evaluate SEC petitions. So these things have been going on simultaneously. That subtask two report we received on November -- or it was dated November 30th, so Board members have had this also for several weeks, and a copy is in the notebook. And I believe copies of all of these are also on the table.

Dr. Makhijani from SC&A is going to give us a presentation now on both the subtask one and subtask two findings of our contractor, so Arjun...

DR. MAKHIJANI: Thank you, Dr. Ziemer. I'll go through the evaluation of NIOSH procedures first. Of course we looked at the rule, 42 CFR 83, and the main procedure's in OCAS PR-004. There are a couple of forms, Form A and Form B, which prospective petitioners would use to file the petitions. We reviewed those, and the procedure makes reference to using OCAS IG-001 and 2, which are the external and internal dose procedures for use in SEC petition evaluation. So those were the basic documents that we reviewed. The OCAS documents were -- 001 and 2

1 -- just mainly we relied on the prior evaluations.

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Let's see, okay. So some strengths were noted for the procedures. There is a logical, stepby-step procedure that is set forth by NIOSH, and it allows NIOSH to divide the proposed class into sub-classes, and that has happened before. It allows, in the more easy cases, when SECs are to be designated for a certain sub-class at least to be early designated, as happened at Mallinckrodt, for instance, provides some useful examples. And then it also provides for a dose reconstruction for non-SEC cancers.

One of the bigger findings was that it doesn't -- the NIOSH procedures currently don't contain detailed guidance on how to calculate maximum dose. I noticed that Dr. Neton made some reference to this earlier in his presentation on Pacific Proving Grounds. This -- this became something of an item in Iowa, to which Dr. Neton also made reference this morning. It's because -- we're calculating a maximum dose with plausible assumptions, and those terms need to be defined in terms of -- not in

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terms of actually putting numbers on a maximum dose, which would vary from site to site and job type to job type and so on, but defining some criteria where there -- where maximum doses might be considered reasonable so that the maximum doses don't become arbitrary. we believe that some -- some guidelines along the lines of actually discussing job categories, evaluation of data integrity, how dose estimation for unmonitored workers might work -- this might be helpful in defining the limits for what might be considered, you know, a scientifically reliable maximum dose rather than an arbitrarily high maximum dose. One of the most important and difficult points was what is the relationship of 42 CFR 83 under which maximum doses are calculated with plausible assumptions to the highest doses under worst-case assumptions under 42 CFR 82. Now under the individual dose rule, 42 CFR 82, the highest doses are not supposed to involve uncertainty. And the way we understand that is when NIOSH declares that a certain number is, under scientifically-reasonable worst-case assumptions, regarded as a maximum dose, that -

- that that is the maximum dose, which is used only for denial.

Under 42 CFR 83, if an SEC petition is denied on the ground that maximum dose with plausible assumptions can be calculated, that dose could be used to compensate people. And the rule is silent and -- and the guidelines are also silent on the question of the relationship between these two kinds of maximum dose. And we believe that because the maximum dose under 42 CFR 83 methods could be used to compensate, as well as deny, that should not wind up being higher than the highest worst-case assumption dose that will involve no uncertainty under 42 CFR 83.

That -- we've suggested that an uncertainty which requires that the worst-case dose under 42 CFR 82 always be higher than the maximum dose under 42 CFR 83 would clarify what is now a problem, at least in theory. And this did come up, indirectly but not in a specific example, at Mallinckrodt where we did raise the issue in a site profile review, that some of these doses had the potential to wind up being higher than calculated worst-case doses for --

for that site.

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This was a theme regarding survivor claimants that came up in our review of interview procedures under 42 CFR 83. Now NIOSH does have provision for working with petitioners quite extensively and helping them prepare the petition so that they qualify after a petition is submitted. But we continue to feel that survivor claimants may have great difficult -who want to become petitioners or -- or survivors who are not claimants who may want to become petitioners, may have a particularly difficult time and -- and some -- to level the playing field, some special assistance does need to be provided for in terms of incidents, in terms of working conditions of the plant, perhaps explaining the site profiles and -- and things like that to prospective petitioners. And -- and you know, post-petition there is assistance, and we haven't evaluated any detailed case study on that, but pre-petition at least some -- some more level playing field would appear to be called for for survivor claimants who want to become petitioners. We listed some other concerns. Currently NIOSH

does provide for a interview with petitioners, but it's not required as part of the guidelines, and we think that a detailed interview with the -- with at least one of the petitioners is very important and should be required as part of the guidelines. It might obviate misunderstandings and difficulties and -- and if -- if an SEC petition is denied, it might make it easier for the petitioner to understand the basis of it if an interview had been conducted, and NIOSH might also receive more clarity as to where the petitioners were coming from in that regard.

There -- there is the -- the guidelines, as well as the rule, do say that the -- some data from the site should be used as a starting point, and provide some of the basis for -- for maximum dose estimation, or dose estimation, with sufficient accuracy. But this is -- how much is adequate, what kind of data can be used from other sites, how it would be married, I think -- I think -- we think that the guidelines could use some examples, at least, based on -- on -- on the work that NIOSH has done so that it's a little more clear. We

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realize that the rule was written

prospectively, but now there's -- there's a

great deal of experience with SEC evaluation,

so many of these things are not necessarily

criticisms of the guidelines as they were

issued when they were issued, but with -- with

some hindsight, perhaps the process might be

smoother if -- if the guidelines did -- did

have some clarification based on the experience

that had been gained.

The last issue related to health endangerment. This came up this morning in regard to the workers who worked less than 250 days. The example that's in the rule and in the guidelines in regard to the external criticality accidents certainly provides an example of where there would be an endangerment in regard to an incident. There's no corresponding example for internal dose. Certainly there have been circumstances at various sites like Mallinckrodt and others where one might imagine doses -- and one can calculate doses based on available data, I believe -- in periods shorter than 250 days that might be comparable in health endangerment

1	to other places where NIOSH has declared health
2	endangered for more than 250 days of work. And
3	I think in the case of internal dose, the
4	guidelines are rather silent, and and there
5	is no corresponding clarification about what
6	the process is for people who are primarily
7	exposed to internal dose and how to proceed for
8	workers who worked for less than 250 days.
9	And then we have some suggestions for
10	improvement that are based primarily on on
11	the findings, which I won't go through. You
12	can look at the list, and they correspond to
13	the findings I've just gone through.
14	So that's the review of the guidelines, so
15	perhaps
16	DR. ZIEMER: Let's pause a minute and see if
17	any of the Board members have questions or
18	comments on this material.
19	(No responses)
20	Okay, let's proceed.
21	UNIDENTIFIED: (Off microphone) Jim Jim
22	might have a comment.
23	DR. ZIEMER: Oh, Jim, please.
24	DR. NETON: Are we ready? I
25	DR ZIEMER. Oh sure veah go ahead

1	DR. NETON: Okay. I just wanted to make a
2	comment on I think it was maybe your first
3	slide, Arjun, or the second one, I'm not sure -
4	- but it has to do with this maximum dose used
5	to compensate or deny. This has been sort of
6	an ongoing misunderstanding or miscommunication
7	I think we've been having, and I'd like to
8	maybe one more time try to address the issue
9	and see if we can come to some some
10	clarification on this.
11	DR. MAKHIJANI: (Off microphone)
12	(Unintelligible)
13	DR. ZIEMER: Slide four four or five maybe.
14	DR. MAKHIJANI: (Off microphone)
15	(Unintelligible)
16	DR. NETON: I don't have the presentation.
17	DR. MAKHIJANI: (Off microphone)
18	(Unintelligible)
19	(Pause)
20	DR. ZIEMER: Go back let's see, strengths
21	there you go.
22	DR. NETON: I hope I can do justice to this,
23	but let me see if I can explain what I think
24	Arjun is saying and then what we our
25	position is. 42 CFR 83 allow for maximum it

establishes the fact that we need to be able to establish a maximum dose with plausible assumptions in order to deny a class. If we can -- if we can establish a maximum dose under plausible circumstances, then the class could be denied. That would be a basis for denying the class. That's the way the regulation reads, at least in our mind.

Now, if that's the case, that does not mean that when we -- if the -- if the class were denied, that we would necessarily use that maximum dose to do dose reconstructions under 42 CFR 82. We're not required to. That would just be a bounding, plausible analysis to demonstrate that yes, we can indeed put some plausible upper limit.

When it comes to doing dose reconstructions, if there is a more refined method available at the time or becomes available, we can refine it and use a better estimate. In some situations, though, it may be that we will have no additional information. The maximum plausible dose that was used to deny the class will be-could become the best estimate. It's no longer at that point a maximum dose. It's the best

estimate that we ever feel that we could come up with.

So I think that might clear up some confusion because it -- they're two totally different concepts. The maximum plausible dose for SEC petition analysis is very different than the maximum dose that we use in the efficiency process to deny cases.

DR. ZIEMER: Jim, I thought that Arjun was suggesting that when you went back and did the actual dose reconstruction you might come up with yet a higher number. Was that the canono.

DR. MAKHIJANI: Dr. Ziemer, the point of -- of this finding is that whatever the terminology, when you go back to 42 CFR 82, after denying a petition, and you've developed a method -- presuming you don't have more information and so on -- and we take the case of Mallinckrodt. A certain plausible dose method was developed, and if that had been denied and you didn't have more information, you would then have proceeded to calculate doses by that method to compensate or deny people, depending on POC.

Now the concern here is that that number -- now

at Mallinckrodt some cases had actually been denied using a different approach that was called highest dose with worst-case assumptions. Now at that point it should not be the doses that are being used to compensate people -- to deny people should have been less than doses that are being used to compensate people. And the point of this comment is that there should be some restriction in going from the SEC rule after denial as to how those cases are handled. And we do recognize that there's a distinction between the terms.

MR. GRIFFON: (Off microphone) (Unintelligible) it's more of a having to go back to those cases and reassess (unintelligible) --

DR. NETON: Well -- well, right, but I mean -- I think what -- this -- this could happen very frequently where we would use worst-case assumptions to deny cases under 42 CFR 82 because it's expeditious and it gives the claimants an answer fairly quickly, while we're continuing to do more research to refine our ability to do dose reconstructions. At the end of the day, if NIOSH finds no more information other than that worst-case assumption and it's

1 a plausible worst-case assumption, it could in 2 effect then become the best estimate for those 3 dose reconstructions and then could be used to 4 compensate cases. It's no longer a worst-case 5 assumption; it's our best estimate. There's sort of a nomenclature issue here, I 6 7 think. Our -- 42 CFR 82 allows us to do best 8 estimates, and if the best estimate is a worst-9 case assumption, it -- that's what it is. 10 it depends on how much research we can 11 accomplish --12 MR. GRIFFON: (Off microphone) (Unintelligible) I think we're really there, from 13 14 (unintelligible). 15 DR. NETON: Okay. 16 MR. GRIFFON: (Off microphone) (Unintelligible) 17 I mean I think, you know, it's the question of 18 if you end up using the maximum plausible dose 19 to actually -- you -- you can't do any better 20 estimate --21 DR. NETON: Right. 22 MR. GRIFFON: -- and you have to apply it, and 23 you look back and you say oh, these previous 24 cases we used a maximum but it wasn't as high 25 as this maximum now, then you have to -- you

1 know, so --2 DR. NETON: Wait, wait, wait --3 MR. GRIFFON: -- I think that's what Arjun's 4 saying, isn't it, that you have to go back and 5 reassess those cases if they were denied based 6 on --7 DR. NETON: A maximum plausible -- a maximum 8 plausible dose under the SEC process is totally 9 different from the maximum plausible dose under 10 42 CFR 82, or could be. 11 MR. GRIFFON: Yeah, yeah. 12 DR. NETON: It could be the same --13 DR. ZIEMER: But it could be the same --14 DR. NETON: -- but (unintelligible). 15 DR. ZIEMER: -- in the case you mentioned. 16 DR. NETON: Yeah, all we have to do is 17 establish that a maximum plausible for SEC 18 petitions is -- can be done. But when we go to 19 do the analysis, we don't -- we would do a best 20 estimate. If the best estimate is equal to the 21 maximum plausible estimate, then it -- it's 22 totally legitimate within the requirements of 23 42 CFR 82 to use that. So I don't see that 24 there is a disconnect. 25 DR. MAKHIJANI: (Off microphone)

(Unintelligible) (on microphone) to clarify the comment. Under 42 CFR 82, as I understand it, there's a certain kind of dose which is to be used only for denial. That's the kind of dose, when you wind up above 50 percent, NIOSH and ORAU actually go back and recalculate a better estimate, if they can. And the worst-case assumptions efficiency procedure is used only for denial.

This finding is directed only at that kind of dose, that that kind of dose, which is -- which it is said in the rule will not involve uncertainty -- should truly be the highest number you could come up with under any dose reconstruction procedure because it's the highest dose that is promised to the public will not involve uncertainty. I don't understand how you can say that we calculated a highest dose that will not involve uncertainty, and then, by another procedure under the same program, come up with another number that's bigger.

DR. NETON: Okay.

DR. MAKHIJANI: That's -- that's the problem
I'm having.

1 DR. ZIEMER: Okay, I underst--2 DR. NETON: I understand. You're actually 3 arguing the reverse, which is if we've denied 4 cases based on maximum plausible, then we 5 should not have a higher maximum plausible for 6 SEC, and I agree with that. I'm sorry for --7 if I was the only dense one in the audience, 8 but I'm glad (unintelligible) --9 DR. ZIEMER: It's kind of a terminology issue, 10 really. 11 DR. NETON: Right. 12 DR. WADE: Arjun's point, while it stands on 13 its own strength intellectually, is also of 14 great value for us to hear what he's saying as 15 it relates to our ability to communicate 16 consistently with people. 17 DR. NETON: I apologi --MR. GRIFFON: 18 Well said. 19 DR. ZIEMER: Thank you. Okay, we'll proceed 20 with the subtask two --21 DR. MELIUS: I -- I have a --DR. ZIEMER: Oh, I'm sorry. Jim, please. 22 23 DR. MELIUS: -- and doing that, and -- first of 24 all, I thought your review of these procedures 25 was -- was helpful and I do sort of want to go

through a couple of the concerns and so forth because I think we're -- I'm not sure we need more (unintelligible) and I think we all are in the process of addressing some of them, and I think we need to acknowledge that. And one is the issue of the 250 days, and I think -- as we discussed, and I think coming out of what we're doing with Pacific Proving Grounds I think we'll have more clarification on how to do deal with that in both -- there's some regula -legal, regulatory -- you know, regulationrelated issues as well as sort of how we -sort of practical way of addressing certain situations where -- where that comes -- may come up and -- and so -- but I do think it -it was helpful.

The one comment I'd like to hear back from either Larry or Jim is this -- it's other -- under the other major concerns slide was the last point, which was no procedures on determining the breadth of the class when NIOSH finds that it cannot reconstruct a claimant's dose. It -- and I've just been trying to understand how you -- I mean I -- I think -- comments that there were no written procedures.

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There's nothing in place yet. I think you're doing it, just -- you know, evaluating these petitions and so forth and just sort of your response is well, should we try to formalize that in some ways that -- as a guideline or -- I -- you --

DR. NETON: Yeah.

DR. MELIUS: -- you know, and to me, the other (unintelligible) corollary to that is this whole issue of how do we divide up a proposed SEC into groups that can be constructed -reconstructed and those that can't, and there's sort of this class definition thing, which I --I also think that you're wrestling with, but -and you've never -- you've sort of accepted -you sort of present this and we -- we never really talk much about the rest of the universe there, the years you're not presenting to us or -- or groups and so forth and -- and --DR. NETON: I agree. For the record, I -- I do agree, there's no procedure for determining the breadth of the class. But I think, operationally, this is -- this is very much driven by the availability of the data. data speak for themselves, and how one would

1 proceduralize that is -- would be difficult for 2 me to envision. I mean we could say exactly 3 that sort of thing. But you know, if you -- if 4 you have a class of workers who worked the 5 third shift and they walked around the plant and had no monitoring data and had similar 6 7 exposures to the people on the second shift, 8 then that sort of just funnels itself right 9 into that definition. The fact of the matter 10 is, we rarely have that type of a situation, 11 and you start looking at classes of workers who 12 may have been all over the plant and you can't 13 necessarily put them in time and space 14 anywhere. I'm not sure how one would -- would 15 narrow it any better, other than a detailed 16 analysis of what we have in our hands at the 17 time. 18 DR. MELIUS: (Off microphone) (Unintelligible) 19 I thought, but now I'm really in trouble 'cause 20 I've invoked legal counsel. 21 MS. HOMOKI-TITUS: No, I -- I just wanted to 22 remind you --23 DR. MELIUS: Or provoked, I guess is... 24 MS. HOMOKI-TITUS: -- NIOSH has also limited 25 what they present to you by what the petition

1 is, so years you're not seeing are because 2 they're years that were not petitioned upon. 3 DR. MELIUS: (Off microphone) Yeah, 4 (unintelligible) --5 MS. HOMOKI-TITUS: So if you get a Y-12 6 petition for ten years, eventually the Board 7 will get an SEC response from NIOSH on those 8 ten years, but not the 20 that followed it. 9 DR. ZIEMER: But we also have cases where NIOSH 10 initiates the class based on their own findings 11 12 MS. HOMOKI-TITUS: Right. 13 DR. ZIEMER: -- and I think this may be --14 MS. HOMOKI-TITUS: (Off microphone) Jim's 15 response (unintelligible) --16 DR. ZIEMER: -- if you find a group, I think 17 the issue is how do you know that you've encompassed the right subset or subsets to --18 19 to define the class. And sort of intuitively you are doing something, and I guess Jim may be 20 21 asking does that need to be codified in some 22 way, or do you just describe what it is you do 23 to find -- I guess that's (off microphone) 24 (unintelligible) --25 DR. NETON: We could certainly attempt to do

that. I mean --

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DR. MELIUS: What I would suggest would -- you know, think of it as a suggestion and -- is that you include a step in this process that, after you've done or -- you know, done a lot of the evaluation, you understand what the critical datasets and so forth, you sort of examine that and say what -- what does this say about who should be included, not included, and are there other groups that the same facts apply to (unintelligible) the same circumstances apply to 'cause why -- 'cause they're going to get -- end up in the SEC class one way or the other. They're -- they're going to end up -- we shouldn't make them have to petition 'cause you're going to identify them when you go to reconstruct their doses, presumably, and we'd have to go through the whole process again. It seems to me it would be -- could be -- it could be more efficient to sort of do it in one step and...

DR. NETON: I think, though, that to a large extent the draft guidelines put together by the working group are going to help with that because we're now -- we're now focused on

looking at not only what was monitored and who was monitored, but what was not monitored. I mean we're required now to go back and look at all these, as John called, exotic -- I'll call ancillary nuclides of exposure and say well, you know, there was also plutonium, americium, curium, and were -- were there monitoring programs in those areas. And then we would define -- it -- it's almost -- ends up being defined then by nuclide type, which is not really helpful, but then you have to figure out what work classes would fall into exposures for those things.

The good thing is right now we're investigating Calutron/Cyclotron operators at Y-12. You know, they were exposed to isotopes other than uranium, and so we're -- the procedures that the workgroup put together are funneling us towards that. I think that'll be helpful.

DR. MELIUS: Yeah, and again, not trying to -- think we should try to make unnecessary work, but the data's all in front of us. We're evaluating it. Let's take advantage of that time when we're discussing it to decide what more needs to be done so that way we don't have

1 to come back to it two years later or whatever. 2 I think that would worth thinking about. 3 DR. NETON: I totally agree. I think, you 4 know, having done this a few times now, we have 5 a lot more knowledge as to -- as to what we're doing and what's available, and --6 7 DR. WADE: And just a comment for the record, 8 this issue of who's to be included and who's 9 not to be included, I think we have to consider 10 inviting into the discussion our colleagues 11 from DOL who often are left with making these 12 decisions once the Board has made a 13 recommendation and the Secretary a 14 recommendation. So I think there is -- there 15 are many voices to be heard for us to do this 16 right. I don't think we started out doing it 17 perfectly right, but that doesn't make us bad people. But I think we can evolve the process 18 19 as we go. 20 DR. ZIEMER: Okay. Okay, thank you. Arjun, if 21 you'll continue, please. 22 (Pause) 23 DR. MAKHIJANI: So this was the second report. 24 This -- this really has two parts. One -- one 25 was based on what had happened with the review

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SEC petitions, what we might suggest as Board procedures, and the second was what we might suggest for our own procedures. This was subtask two, which you also commissioned. For the Board procedures, we suggested that we think about it in three phases, and this is I think fairly -- has a fair degree of overlap with what the Board working group did, but it sets it forth in kind of a time frame that may be a little bit different. The phase one would be preliminary steps taken immediately after NIOSH qualifies a petition for evaluation. phase two would be during the time that NIOSH is actually evaluating the petition. And phase three would be after NIOSH has submitted an evaluation to the Board for consideration. Okay. So the most important part of phase one was that we thought that it might be helpful if NIOSH would start with some implicit evaluation plan when it begins evaluating the petition. mean it's got a certain amount of data there in front of it, documents, perhaps dose reconstructions that are relevant. And if NIOSH could submit a plan, you know, to present the documents and how it is proceeding to

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evaluate the petition at about the time it begins to evaluate itself the petition, the Board, through a working group, could begin that process to identify the issues. And that would make the process of evaluation of the petition involve the Board more closely, rather than having a completed product presented to the Board after the evaluation is complete. And doing that review by the Board serially and -- that would cut down the time and also there'd be a greater likelihood that all of the relevant issues would be put on the table during the process of NIOSH evaluation of its petition. But that of course depends -- the key step is that when NIOSH begins its own evaluation that a relatively rich plan that it itself is following, together with the documentation, is -- is supplied to the Board. So phase two -- this derives a lot from -- from the experience that we had with Mallinckrodt which went on for quite a long time, and -- and partly is suggested to compress that procedure. The dose -- example dose reconstructions that would illustrate the issues would be defined and provided. NIOSH would do them itself. The

Board would review the material. And the overall goal of this step would be to ensure --would be focused on these example dose reconstructions to ensure that the issues that are relevant to the feasibility of dose reconstruction would -- with sufficient accuracy would be examined in depth so that NIOSH, as well as the Board, could -- could arrive at a conclusion whether -- as to -- as to the feasibility.

Then in phase three, the phase three is relatively straightforward. It's spelled out in 42 CFR 83 quite well, and -- and the Board has already been following this: to hear from NIOSH, to hear from the petitioners, to hear from others and consider all the different points of view and decide whether a further review is needed or whether you can take action. So the phase three part of it is -- is not at all new, whereas the other two phases would be basically drawn on the experience of what the Board has done in the past year to review SEC petitions.

So that -- I don't know if you want me to pause here or just go on through to the end. We also

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suggested draft procedures for the Board contractor, and they're spelled out in quite a bit of detail, but basically it would be to review the petition and associated documents and to determine what kinds of partial dose reconstructions would be needed in order to clarify the issues regarding feasibility of dose reconstruction. We propose to interview site experts and at least one of the petitioners as a required part of the evaluation process. If there is a full -these steps involve the evaluation after NIOSH has submitted an -- its own SEC petition evaluation, if SC&A or the contractor is asked to review the whole evaluation that NIOSH has done, this would be -- it would be a rather lengthy and detailed process that would involve these interviews that would involve looking at dose reconstructions and of course working under the direction of the Board and the working group to assess the representativeness of the data, the adequacy, validity and to see whether the technical issues that had been raised in regard to feasibility have been satisfactorily assessed.

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Now the Board did ask us to examine what we might do when there is a site profile and a site profile review and those minimum things are -- are mentioned here. When there's -when there's no site profile review, we would not propose to undertake a site profile review as part of an SEC petition process. It would be more targeted review of the site profile that was relevant only for those issues in regard to maximum dose reconstruction. I think this has been exemplified by some of our presentations yesterday and today where we did make an effort to sort out those issues -while they may be significant for dose reconstruction -- but we thought can be resolved. Like, for instance, the geometry -the geometry of the workplace in relation to the badge when it is known. This is -- this is something that can be done and doesn't rise to the level of whether dose reconstruction is feasible, but merely how it should be done. Whereas there were other issues that have been raised in regard to Rocky Flats and Y-12 that would need to be resolved to determine the feasibility. So we would sort those out and

1 focus only on those issues where feasibility 2 was involved and nothing else. Of course we 3 would not be dealing with minimum or -- or best 4 case doses at that stage, but just the 5 feasibility. And similarly when there's no site profile, 6 7 then we would perform a focused review of the 8 conditions at the site -- radionuclides, job 9 types and so on -- with the same targeted idea, 10 that what is relevant for maximum dose 11 reconstruction under plausible assumptions for 12 that site. 13 There's a little bit of a complication that --14 it's not really discussed at length, in the sense that these procedures don't cover what 15 16 might happen in regard to drawing data from 17 other sites. We've been silent on that. NIOSH 18 does use that, and we would just follow in our 19 evaluations what NIOSH had done, not --20 presumably not be initiating our own 21 investigations on other sites. 22 In regard to partial reviews, there is 23 provision in the task order for partial 24 reviews. I imagine that that would happen in 25 what I described as phase two of the Board

1 procedure. Should the Board choose to ask for 2 some support from your contractor, then you 3 would define the issues during the process of 4 NIOSH evaluation of the petition and we would 5 work along with you on those issues, as we did 6 at Mallinckrodt and Iowa, and as we are doing 7 in -- so far in the case of Y-12 and Rocky 8 Flats. 9 That simply gives you the review preparation 10 team. 11 DR. WADE: Just to expand on what Arjun said, 12 part of the task we put in place with SC&A 13 allows for the Board to task SC&A with assisting it in a petition evaluation review. 14 15 And Arjun talked about the two types, sort of a full review and then this -- this task-16 17 specific. So -- I mean all of the mechanisms 18 are available to you now to use as you might 19 like, and I think Arjun made that fairly clear 20 in his comments. 21 DR. ZIEMER: Okay, questions or comments, Board members, on this -- okay, Dr. Melius. 22 23 DR. MELIUS: Yes, I would just offer up that I 24 think that, given the time frame for trying to 25 address these, that -- that I think we possibly

1 want to avoid the complete reviews, that --2 that -- you know, the partial review -- and I -3 - I hate to call it partial, I'd rather call it 4 targeted or something that doesn't 5 (unintelligible) --UNIDENTIFIED: (Off microphone) 6 7 (Unintelligible) focus (unintelligible). 8 DR. MELIUS: -- for the focused review that we 9 -- would be more appropriate because --10 particularly where there's been a site profile 11 review. I think when there's no site profile 12 review, then I think this becomes lengthier just -- you know, particularly for a large 13 14 site. Now for a small site or something, or --15 not a -- a less complicated site, that may be 16 different. But for a large site, then without 17 a site review it -- it's going to take some time to -- to make it -- make it through the 18 19 process. 20 And this is a question both for -- I guess for 21 -- sort of the suggestion for the Board, but 22 also for -- for NIOSH that when we talked 23 yesterday about the -- a workgroup report, we 24 made the suggestion that there be -- that 25 currently NIOSH develops its evaluation plan

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very early and it's a very general plan, so this leads to the general steps that we're going to take. Seems to me that there's some point, after NIOSH has had a -- maybe a little bit more of a chance to review the materials, think -- think about the petition, so forth, that there -- there would be a time when we could pull together -- you know, that NIOSH would have more specific plan, what were going to be the crit-- what they saw to be the critical datasets and -- and so forth, that that would be the time to put together -- pull together NIOSH, a workgroup and the contractor, sort of sit down -- particularly the contractor staff would have been involved in the site -site profile review, to sit down and -- and sort of look at how do we -- you know, where do we go from here, how do we go through and review the -- the portions of that site profile, the portions of the data that are going to be important for the SEC evaluation. And I -- if I understand the task order so forth -- process right, I think that would be an appropriate way of -- of proceeding. The -the issue would be scheduling that first

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meeting, but from there there should be -- then be sort of a work plan and everyone be able to sort of schedule out what would be involved. I think when I was listening this -- when we were discussing the Y-12 site profile review, in some sense that's already being done there. I mean I think -- gives a pretty good idea what the critical parts are, you know, that need -and I think even for Rocky Flats, I thought I heard -- heard the same and so forth. but to me, that would be a way of having a procedure in place, avoiding a very lengthy step-wise -- you know, serial process, but let's try to do some of this in parallel. Keep the -- our contractor's task sort of focused in relationship to the -- to the SEC review. Is -- is that making sense, I guess is my question to my fellow Board members and NIOSH? DR. ZIEMER: Well, in a sense what we're doing now is -- is targeted reviews of the site profile to assist us in -- in the SEC evaluation, so that's -- in practice is what's

MR. GRIFFON: I -- I was going to -- I mean as I read this section, I was thinking the full

1 review is actually what we're now calling the 2 targeted site profile review 'cause there --3 it's a full review of the -- of the issues that 4 may affect the SEC, you know. That's -- that's 5 sort of how I was interpreting it. But -- I 6 mean I -- if you want to modify the language, I 7 think that was the intent. DR. MELIUS: Yeah, it's a little bit mixed in 8 9 how it's presented, 'cause I think we're also 10 going back to sort of Mallinckrodt examples and 11 so forth where we got -- we were really using 12 the site profile review --13 MR. GRIFFON: Right. 14 DR. MELIUS: -- in lieu of a -- a SEC review, 15 and I think that has some disadvantages in 16 terms of -- of some of the questions it can 17 focus on, and that was some of our -- our 18 problem. We weren't necessarily getting --19 weren't having our contractor, you know, answer 20 some of the key questions or give us advice on 21 some of the key questions. 22 DR. ZIEMER: All right. We'll hear from Larry 23 and then from John. 24 MR. ELLIOTT: You know -- yes, we have been 25 dwelling on the early history of our

1 development of the evaluation reports on 2 petitions, and I would say that our 3 Mallinckrodt experience and our Iowa experience 4 are -- are learning experiences, one we're 5 building from. The plan that we gave you -- that we give you 6 7 on every evaluation of a petition is a generic 8 plan. It is not specific in detail. It is, we 9 hope, comprehensive in its general--10 generality. What's lost -- in my understanding 11 of the proposal that has been made by the 12 working group and what I see in SC&A's comments 13 in their review about another plan, a more 14 detailed plan -- is that we're struggling with 15 our -- we're struggling really hard trying to 16 come forward in 180 days with a scientific 17 basis to make a recommendation to add or deny a 18 class. And I'm not sure -- I can't -- I can't 19 seem to grasp how we would do that in 20 conjunction of providing a detailed plan along 21 that trail. 22 I see it more as what I can -- what I can 23 understand in my mind is a more concerted coordination effort where we sit with the 24 25 working group and SC&A and say here's where

1 we're at, here's the -- here's the -- the 2 salient issues that we're wrestling with on how 3 to evaluate this petition. And you know, I 4 think if we can -- if we can approach it as a 5 more -- a better coordination effort, we're better off. I hate to see -- to me, a plan, an 6 7 additional detailed plan within the context of 8 a 180-day time requirement, is make-work for 9 And I'm not sure that's going to get us a 10 lot farther down the road or a lot quicker down 11 the road. But I do believe that a better, more 12 concerted coordination effort can get us to 13 where we want to be. 14 DR. ZIEMER: Your concern then is that you may 15 spend an excessive amount of time on the plan 16 versus actually doing the job that needs to be 17 done. 18 MR. ELLIOTT: Yeah, we don't have a lot of time 19 20 DR. ZIEMER: Yeah, right. 21 MR. ELLIOTT: -- in 180 days to create another 22 plan. 23 DR. ZIEMER: No. 24 MR. ELLIOTT: And that's why you haven't seen us come forward with a different version of a

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1 plan for each SEC petition. 2 DR. ZIEMER: Yeah. 3 MR. ELLIOTT: We gave you a general -- what we 4 consider --5 DR. ZIEMER: A generic approach, yeah. 6 MR. ELLIOTT: -- a generic, comprehensive plan 7 that spoke to these are the types of things 8 that we would have to consider in evaluating a 9 petition. And we've stopped short of providing 10 any further detail on that. I certainly want 11 to work, you know, in a more coordinated 12 fashion with the working group and the Board 13 and SC&A on getting to the end product here. 14 But I'm not sure that -- in my opinion, I don't 15 think we can just generate another plan and 16 it's going to help us get there. 17 DR. ZIEMER: Thank you. John. 18 DR. MAURO: I'd like to bring up a practical 19 issue related to managing budgets and how we're 20 proceeding with our work. When we have a site 21 profile such as Y-12 where we perform our 22 review, deliver our product, the bottom line is 23 that -- the way we try to manage it is we try 24 to deliver that product to you -- thick book --

within 1,000 work hours. That's how we manage

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ourselves. We set aside 150 work hours for the closeout process. Okay? So let's say we're dealing with a site profile that is -- does not have SEC implications. That plan works. I think it's very doable to, within 150 additional work hours, to engage in the dialogue, hold our meetings with the working groups, close out issues, put it to bed -- if we can.

However, what I am concerned about is -- let's use now Y-12 as an example. Okay? We've delivered our Y-12 work product. We did -delivered it within budget. But right now I have in the bank 150 work hours to support the closeout process. Well, the closeout process, as we all know, is now really part of the SEC process. I don't know what we will be called upon to do. Previously Mark had mentioned that we had these 6,000 records. Probab-- there might be some things that you may call upon us to do. It may simply be on the back end to take a look at whatever the results are. You may call upon us earlier to be more involved. What I -- I guess where I'm going with this is that keep in mind that when we're in an SEC

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mode, we are going to run into back-end work hour problems; 150 work hours will blink out of existence like that (indicating) when we get involved aggressively in the SEC closeout process.

DR. ZIEMER: Thank you. Okay. Arjun? DR. MAKHIJANI: Yeah, when we were drafting the -- the report, both for Board procedures and SC&A procedures, we did think it useful to kind of focus -- and maybe it's my lack of understanding of, you know, institutionally what's involved in developing a site-specific plan. And perhaps the words that Larry used would be -- would be more appropriate. And I think if that's what we've been doing now in regard to SEC and focusing those issues, that -- that would be appropriate. I -- I don't know, but we -- certainly some site-specific information from NIOSH is necessary as to how -- where NIOSH is and how they're evaluating in the process.

And -- and the one point I think where a lot of effort is needed on SECs where no effort, or very -- well, I wouldn't -- I shouldn't say no effort, but some internal effort is put in when

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we prepare site profile reviews and in coordinating with the folks that are doing dose reconstruction audits to figure out what's going on in individual dose reconstructions and so on, but while we inform ourselves during the review, there isn't a huge effort around dose reconstructions when we do site profile There's some, but there's not a lot. And it seems to me, the way it has worked out and what seems to best exemplify feasibility under SEC, that a lot of effort has to be devoted in developing, defining and then having some rubber meets the road test around are these methods actually applicable in real dose reconstructions. And I think that is -- that's the big point of departure between just doing a site profile review, other than documentation and verification and so on with various other...

The other -- the other big difference is -- in site profile reviews, just to remind you of what our own procedure has been and what we've been delivering to you, is we don't cover all issues. We highlight issues -- we call them vertical issues. We don't promise you that

1 we've chased every last thing down. 2 when you're doing an SEC petition review, 3 within the framework of a more limited dose 4 reconstruction idea, you do have to chase down 5 the last thing on feasibility for the last answer for all cancers and for all members of 6 7 the class. And so the goal is very different. 8 And while we've been kind of muddling along in 9 terms of -- if I might use that phrase, you 10 might excuse me -- that to -- to move ahead 11 with site profile reviews, I do think, having -12 - having spent quite a lot of time drafting these reports, that there is a great deal of 13 14 merit in having something that is focused on 15 SEC reviews. 16 DR. MELIUS: Can I respond to a couple of those 17 points? 18 DR. ZIEMER: Yeah, let me have Mark -- Mark was 19 up (unintelligible)... 20 MR. GRIFFON: (Off microphone) (Unintelligible) 21 I mean I was just going to say that I think what -- Larry's comment I think speaks to the 22 23 spirit or the goal of our intent, which was 24 earlier involvement with the Board and the 25 contractor -- and our contractor and, you know,

1 I don't know that -- I -- I tend to think --2 you know, we don't need a -- my fear here, I 3 think, is that if -- if we put a step in to 4 develop a plan -- as long as we have a 5 commitment for earlier involvement, I'm not sure we needed a commitment for a plan because 6 I think that plan may be out the window by the 7 8 end of the 180 days, anyway. So you know, I 9 think that the -- the real thing we need is 10 that -- that -- that commitment for the earlier 11 involvement and -- and get together, focus, 12 find out what -- you know, put the issues on the table, the likely issues that are going to 13 14 come up in this SEC review, and then go forward 15 from there, you know, so --16 DR. MAKHIJANI: Yeah, Dr. (unintelligible) you 17 know, I haven't consulted with my colleagues, 18 obviously, but I -- I do take -- I do take the 19 spirit of Larry's comments, and I entir--20 entirely agree with you. There's -- there's 21 not a bureaucratic intent in -- in what we 22 propose, but --23 DR. ZIEMER: Right, understood. 24 DR. MELIUS: Yeah, first of all, to answer 25 John's question, yes, you will get -- you will

get paid, that's why we (unintelligible) the task for the SEC reviews --

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DR. ZIEMER: Notice who's guaranteeing it, though.

DR. MELIUS: -- we just want to make sure you don't get paid too much, so we want to focus it a little bit, so -- that. (Unintelligible) I thought about this a little bit and yeah, we don't want to (unintelligible) but there are some things that I think you -- from the time you produce your generic plan to the time that you really get going on the (unintelligible) plan, you do, and I think one of the steps that you do -- that you have access to that we don't are the completed site -- the dose reconstructions or those underway for that particular site. And I think you can -- you pull those out, you understand what -- what ex-- what datasets you've used to do those, what have been the problems and so forth. think bringing that experience -- and I think that takes you some time to do, I -- I think bringing that experience forward (unintelligible) not preparing a report but being ready to talk about that with a workgroup

from the Board, with the Board's -- our contractor there, along with whatever has been done to review the site profile and so forth is going to -- you know, that's the plan. That's where you're going to -- we'll develop a plan. Is that plan going to be perfect? No. Does it have to be a written thing that gets reviewed? No. But -- but I think -- you bring that. contractor would bring, you know, the site profile review that's been done, or at least -and maybe -- wherever that is. It may be underway, it may not be completed yet. They would bring that, and from that, hopefully we would develop, you know, a way forward that would be sort of more focused. What our guidelines -- our workgroup guidelines

What our guidelines -- our workgroup guidelines and so -- and the criteria and so forth that we talked about yesterday and talk about more today, I -- I think, again, provide sort of a framework for providing that focus, but I don't think anyone's expecting that, yeah, it's these three datasets that we're going to spend -- you know, and, you know, do this, that and that. It's not that kind of a plan, but -- but at least there'd be a way forward and yeah --

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yeah, we need to do this and meet again in, you know, a month and we'll provide this to you, something like that. And I think the -estimating that it's going to be -- claiming it's going to be done in 180 days or 120 days or whatever, you're not going to know at that point in time. You're not going to know until you figure out, but at least you'd have some sense of what the -- the overall scope and resources necessary and what you'd have to do. But -- but you already have some inform-- you don't -- you'll have information that nobody else has access to at that point in time. MR. ELLIOTT: It's better to bring you all along as we go through it, I agree. I see the merit and the benefit to that. And as Dr. Neton answered your question earlier about our -- our attitude and our feelings toward the -the sufficient accuracy paper that you talked about earlier, we are behaving that way. taking that seriously and we're -- we're modifying how we go about doing our evaluation of a petition, recognizing that the Board wants to see example dose reconstructions if we're saying to deny the class. How can we go about

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doing a dose reconstruction in order to support that recommendation. I think that we would all be better served if we could map out, in a time-line fashion, you know, what has to happen over the course of this 180-day mark and -- and know that there's certain suspense dates in that time line that certain things have to happen. We have to have a meeting with the working group, we have to share with the working group where we're at currently, what -what we're wrestling with. We should vet draft examples and make sure that they're going to be compelling enough, or are we going to have to go back to the drawing board and find some more, give you a better sense of all of the dose reconstructions that have been completed to date and how they were completed and why we think they were done with sufficient accuracy. All of that I think could be mapped out in a time line which may serve us well -- you know, better than a -- creating a plan and then coming back and saying well, that plan didn't work. We've got to create another plan. Yeah. Good. Other comments or DR. ZIEMER: questions?

DR. WADE: I have a comment.

DR. ZIEMER: Yes, Lew.

DR. WADE: I mean I think this has been a very useful discussion and -- and I think the spirit of what Larry put on the table I think is exactly what Arjun came suggesting. So I think while it might have been different terminology, I think there's a great deal of coincidence there.

What I would point out to the Board, though, is that -- and I've given you this piece of paper which speaks to sort of the status of -- of SEC petitions. And I'm not saying you need to do it now, but if you want to think about having that meeting that Larry sort of offered that Arjun is asking for early in the process to sort of sort out these issues, your -- your target of opportunity is limited in that -- I mean if you'll look at this page, you have -now we -- we've -- we're working on Pacific Proving Grounds, Y-12 and Rocky Flats. You can do more on those if you wish. But the only three other qualified petitions are listed there. You might want to choose one of those to use as the first example of trying to put

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1 this process into place, and that is have this 2 early interaction between SC&A, NIOSH and the 3 Board to try and work these issues, whatever 4 you call it. I'm not suggesting you do that, 5 but there's an opportunity to do that. 6 targets of opportunity are somewhat limited. 7 In NIOSH-speak, it's not a petition until it's 8 qualified. 9 DR. ZIEMER: Thank you. Other comments now? 10 Larry, uh-huh. 11 MR. ELLIOTT: The public -- I don't know if the 12 public realizes this, but the document you're talking about is on the table. 13 14 DR. WADE: Okay. 15 MR. ELLIOTT: And I just think that it would be of -- in good order to read, for the record, 16 17 what site petitions we are currently evaluating 18 so that --19 DR. ZIEMER: Go ahead, Larry. 20 MR. ELLIOTT: -- you know, it can be codified 21 there. We have the -- the Pacific Proving 22 Ground, which you have attended to today. We 23 have Y-12, that's petition 28 and that covers 24 the plumbers, pipe-fitters and steamfitters 25 from '44 to '57 and we talked about that also

at this meeting. We also have the Rocky Flats petition, and of course we're way behind on it and we want to get the site profile resolved in order to make the evaluation report stand on its own merit.

We also have the Oak Ridge Institute for Nuclear Studies, and here's an interesting one because almost all of the ORAU folks are conflicted on that and so my folks are doing that, and that presents us another resourcelimiting problem.

We also have Ames, Iowa, which is a very interesting site and my folks are telling me that they're coming along fine with working on that. There's a lot of good documentation that the petitioners have provided us and, you know, maybe that's one you might want to pick up and we -- we start working a time line against.

We also have Chapman Valve that's been qualified, and so we are preparing to -- you know, initiate our evaluation report on that.

DR. MELIUS: On which of -- of those latter three, which ones have site profiles? Or are included in site profiles, that may be...

MR. ELLIOTT: Jim, I would ask you to correct

1	me if I'm wrong, but I don't believe Ames
2	Laboratory has a site profile. Chapman Valve
3	does.
4	DR. MELIUS: That's what I thought.
5	MR. ELLIOTT: Oak Ridge Institute for Nuclear
6	Studies does not. So there's different
7	that's good to know the flavors that we have to
8	deal with here.
9	DR. MELIUS: And on Chapman Valve, is the
10	are you reviewing that, John?
11	DR. MAURO: No, we we have cases we have
12	cases that involve that, and as a result of
13	that we read and review the site profile that's
14	that's relevant, but we do not have we
15	have not been authorized to review Chapman
16	Valve as a site profile.
17	DR. ZIEMER: Okay.
18	MR. ELLIOTT: Also on this
19	DR. ZIEMER: Larry.
20	MR. ELLIOTT: second page I apologize,
21	this was a FAX that we had sent in to us today,
22	and the second page lists a summary of active
23	SEC submissions and petitions, some of which
24	are duplicated from the first page, but some of
25	which here on the second page have not yet

1 qualified. And I apologize for the two black 2 lines here. I'm sorry, I can't -- I can read the bottom line, that's Linde Ceramics, and the 3 4 class has been added and that's been treated. 5 I don't know what -- I think the other top line that's black --6 7 UNIDENTIFIED: (Off microphone) 8 (Unintelligible) 9 MR. ELLIOTT: Yeah, that's NBS -- that's NBS, 10 and so that's been dealt with, as well. 11 DR. WADE: So just to give the Board some --12 DR. MELIUS: I thought they were redacted. I 13 wasn't even going to try, that's the reason 14 I... 15 DR. ZIEMER: Lew. 16 DR. WADE: Well, but just again now, if the 17 Board is -- it wants to in earnest move forward with trying to utilize its contractor, this is 18 19 the universe. Maybe it'd be worth choosing one 20 and having an early meeting. 21 DR. MELIUS: Can I work -- as a question, going backwards. What about Y-12 and Rocky Flats, 22 23 where are we in terms of -- obviously we -- I 24 don't think we can necessarily go through the 25 whole plan as we've talked about, or the

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process, but we do have those that are going to come up shortly -- shortly for consideration. They are, you know, fairly technically complicated. We have site profile reviews underway, but I thought I heard -- what John was saying is that he's sort of running out of room to do -- do much, I mean -- which I think is a legitimate concern on those, and I would think that we'd want to consider invoking our contractor's assistance on both of those, also, in terms of the SEC evaluation portion of it. DR. ZIEMER: I'm going to suggest that you keep that thought in mind, and we'll return to it. Now what I'd like to do is get our document on the board, and then having the input from the two presentations by SC&A and the discussion that we heard and our own document, you'll have a chance to mull this over tonight --DR. WADE: We have time on the agenda tomorrow. DR. ZIEMER: -- and tomorrow we have time on the agenda to move forward with a specific plan and, if so desired, to identify some -- one or more sites. I would prefer that we not try to identify sites until we get through the framework on which we will proceed.

PROCEDURES FOR BOARD EVALUATION OF SEC PETITIONS DISCUSSON PART II (INCLUDING DISCUSSION OF THE PENDING Y-12 PETITION AND A DISCUSSION OF THE SC&A SEC TASK) DR. JAMES MELIUS, ABRWH

So with that in mind, we come to the item called procedures for Board evaluation of SEC petitions. Yesterday Dr. Melius gave us an overview of the document. We had -- and the Board has had the document in advance to look at. I think now is the opportunity to have further discussion on that document, input. And we do not have to necessarily approve the document today, but we might -- we might identify -- if there are any substantive changes to make, identify what those are. Lew, I think you expressed to us the interest of the Secretary's office in the document as it's developed. Is -- is there an interest in having input from them --

DR. WADE: Yes.

DR. ZIEMER: -- as well, let me ask you --

DR. WADE: Yeah, let me -- let me speak to this just briefly. As I mentioned yesterday, the -- I briefed the Secretary's staff on the agenda, and there was particular interest in this item, as you can imagine, because the Secretary is -- is the recipient of your recommendations. So

as I mentioned yesterday, the Secretary is very interested in your giving a full vetting of the document, as you -- as you go through. The Secretary would also like the opportunity to comment on your procedures before you finalize them, and I think we should afford the Secretary that opportunity.

It doesn't mean you can't be guided by what you do and we can't make decisions, and we need to make decisions as we move forward. But I think there is an interest on the Secretary's part to -- to provide you with comment as you move to finalize.

The other thing that I would propose that we do is we need to hear from counsel as to what timeliness means in the context of the law and the rule. And whether we do that tomorrow morning or whether we do that now, I think it would inform the discussion if we had a sense of what timeliness meant.

DR. ZIEMER: Yeah. Well, let's -- let's get the document on the floor and I'll ask Dr. Melius to kick off the discussion here again, and then have the opportunity for Board members to in-put. Go ahead, Jim.

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DR. MELIUS: I think the -- well, I guess first of all, the report. I've heard no comments from other Board members since yesterday when -- when we discussed it. There -- I think Mr. Miller last night in his public comments, or late yesterday afternoon I think, raised -raised two issues, and I have discussed some of those issues with other Board members, that -one was the -- I -- the feasibility, sort of timeliness issue, which I think, you know, could be addressed in this. I think there's the issue of the regulations, as well as maybe your legal interp -- 'terpretations, so that might be better dealt with after we've had this legal presentation tomorrow or whenever. The other issue which is the issue of the use of data from other sites as a -- for a site profile. I guess Bethlehem would be an example, but it's come up elsewhere, and should we develop criteria guiding the use -- use of such data, so forth. And actually in some of my discussions other Board members have mentioned they thought that was something that might be appropriate for us to -- to incorporate in some way. I would actually

1 think that that would require some sort of 2 further discussion, either here or possibly, 3 you know, reconvening our workgroup to work on 4 developing criteria 'cause it's not a -- it's 5 not straightforward to do. I think there's some -- some pitfalls we need to -- to be aware 6 7 of in doing that -- doing that. So I think 8 that's where I stand now in terms of comments. 9 DR. ZIEMER: On the timeliness issue, right now 10 the document speaks to it more conceptually in 11 terms of the Board's interest in moving forward 12 in a timely fashion. In fact, we use -- I 13 think intentionally -- rather fuzzy terms. Wе 14 don't have any statements that say we must act 15 within a certain time frame. On the other 16 hand, it's clear that we do not want this to 17 stretch out. 18 Now we can be informed in terms of any legal 19 aspects of what constitutes timeliness, or 20 maybe not. The attorneys may not want to touch 21 that a bit. So --MS. HOMOKI-TITUS: No, the only thing I can 22 23 tell you is --24 DR. ZIEMER: -- can you give us a timely answer 25 to this question?

1 MS. HOMOKI-TITUS: The only thing I can tell 2 you is that I can share with you how timeliness 3 is used in the law. I can share with you how 4 it's used in the regulation. But to give you a 5 legal interpretation of it, just having been asked yesterday, I cannot --6 7 DR. ZIEMER: Yes, I --8 MS. HOMOKI-TITUS: -- we call them 9 (unintelligible) legal opinions and we don't 10 give them. 11 DR. ZIEMER: Right. Right, thank you. In any 12 event, I think the -- I think the document 13 could stand, in terms of philosophical view of timeliness, without having the legal framework, 14 15 probably. But the other issue on the use of 16 data from other sites basically didn't get 17 addressed in this document. And perhaps that's 18 an add-on that we might want to include and 19 have the workgroup actually deal with that. 20 DR. MELIUS: Can I comment on the timeliness 21 issue? 22 DR. ZIEMER: You bet. 23 DR. MELIUS: Since we won't have a legal definition of -- of it. And this is just a 24 25 thought, maybe. I'm not even sure I'm

1 suggesting we do it, but -- but one 2 consideration should be should we make some of 3 the language more specific in terms of what are 4 some of the situations that may arise that -that would affect timeliness, and what -- what 5 6 are justifiable reasons for (unintelligible), 7 how -- how would we handle procedurally things 8 that the -- for example --9 DR. ZIEMER: Are you thinking of cases --10 DR. MELIUS: -- data -- access to --11 DR. ZIEMER: -- or examples? 12 DR. MELIUS: Yeah, access to a dataset is not 13 possible, critical dataset that we need for 14 doing dose reconstruction is -- or is -- to --15 in -- for evaluating for SEC purposes. That's 16 just not available. To me, that's a very 17 justifiable reason for, you know -- you know, postponing action and so forth -- and that, and 18 19 something like that. And so it would be along 20 those lines sort of how -- how we would deal 21 with tho -- those types of issues. 22 DR. ZIEMER: But again, you're talking about 23 providing examples of how one might --24 DR. MELIUS: Examples as opposed to -- yeah. 25 Once again, I (unintelligible) --

DR. ZIEMER: As opposed to -- (unintelligible),
okay.

MS. HOMOKI-TITUS: You haven't provoked counsel. That was actually going to be an example I was going to give, because the SEC rule does speak to that specific situation.

DR. WADE: Maybe you -- maybe you can do your li-- can you run through your list for us?

MS. HOMOKI-TITUS: Oh, sure.

DR. WADE: I think it'd be --

MS. HOMOKI-TITUS: What I pulled together was out of the actual statute, timeliness is only
mentioned once, and it's under the purpose of
the program, to compensate -- it's a
compensation program to provide timely, uniform
and adequate compensation to covered employees.
Lew also asked me to pull the two new deadlines
that were added -- I think you all probably
know them by heart, the 180 days to make a
recommendation, and then the President -- which
has been designated to HHS -- has 30 days to
provide a determination to Congress, once this
Board provides a positive determination to the
Secretary. And then I was also going to let
you know that the SEC rule, as it currently

stands, and the recommended rule still includes this part, gives the Director of OCAS the ability to determine that records are not available in the timely manner. And this will be considered the same as a determination that the records are not available at all. So those are kind of -- those are the legal places where it's actually included. And then if anyone's really that interested, there's a lot of preamble language discussing it that I can give you copies of.

DR. ZIEMER: Okay. Thank you. Wanda and then Jim and then Henry.

MS. MUNN: Both of the items that are being discussed here are resource-limited. It would be very nice if we had all the time that we needed and if we had all of the resources necessary to do all of the comparisons that we need do -- to do, and nobody was upset about not having their claims or their SECs moved forward last week. But that doesn't apply either to this Board nor to NIOSH nor to our contractor. We're all resource-limited. The more specific we become, the less fluid the process can be. And the less fluid the process

1 can be, in many cases, the more -- the less 2 timely it is likely to be. 3 The document we have before us is pretty well-4 written, and allows the reader to understand 5 that everyone involved would like to see timeliness, would like to avoid the use of 6 7 surrogate sites, but when timeliness is an 8 issue and resources are an issue, we must make 9 hard choices. My personal choice would be to 10 accept the document the way it is. 11 interest of timeliness, I would be more than 12 willing to move it when the other Board members 13 are ready for that. 14 DR. ZIEMER: Thank you. Jim, did you have 15 another comment? 16 DR. MELIUS: Yeah, I would just -- I actually 17 was just thinking, maybe I should just stop 18 here since I've got Wanda -- unusual situation 19 -- agreeing with me on it. 20 MR. GRIFFON: Yeah, I was just about to say, 21 I'm willing to second it if (unintelligible). 22 DR. ZIEMER: Okay, we'll hear from --23 DR. MELIUS: But, however --24 DR. ZIEMER: However. 25 DR. MELIUS: I'm not taking issue, but just a

clarification, I would think it could be helpful that if we took a little bit of time and a little bit of resources among our Board to try to come up with some criteria for where -- how to use data from other sites. I think that's --

DR. ZIEMER: Okay. Thank you.

DR. MELIUS: -- one that would be -- be useful and would hopefully save us some time going in the future.

DR. ZIEMER: All right. Henry Anderson.

DR. ANDERSON: I was only going to address a different timeliness issue as -- as the timeliness here is it seems to be on the front end, the timeliness of getting the SEC petition processed. I was just going to raise the issue which came up in some of our other discussions that if the determination is made that it is feasible -- in other words, you're going to deny the petition -- there needs to be some timeliness in when will these petition -- or the cases be dose reconstructed. I think one of the issues was well, we can do it, but it -- I mean that's why we asked for examples -- but they haven't really done it yet. And as we've

1	heard this morning, there's some people that
2	for the Proving Grounds filed four years ago
3	and clearly somebody looked at them at some
4	time and said we can't do these, and then they
5	just sat.
6	DR. ZIEMER: Uh-huh.
7	DR. ANDERSON: And so if a determination's made
8	it can be done, it has to move
9	DR. ZIEMER: Right.
10	DR. ANDERSON: and somewhere we just need to
11	recognize that there needs to be a timeliness -
12	-
13	DR. ZIEMER: Right.
14	DR. ANDERSON: of being able to complete the
15	cases that are waiting.
16	DR. MELIUS: I think we actually
17	DR. ZIEMER: Other comments?
18	DR. MELIUS: we do have that on page 8,
19	so
20	UNIDENTIFIED: Oh, you do? Fine.
21	DR. MELIUS: Yeah.
22	DR. ZIEMER: Now also I would point out that we
23	could, in principle, accept the document with
24	the understanding that it would be expanded as
25	needed, with additional criteria as they're

1 identified. What I'm going to suggest --2 Well, I want to ask this question, Board 3 members. Is there anything in the presentation 4 by SC&A on what they identified as the 5 procedures for a Board SCA (sic) review, anything that you see there that would cause 6 7 you to want to modify the workgroup document? 8 To a large extent, it's a supplement to it, and 9 more of a procedural supplement, but I do want 10 to provide that opportunity if you think any 11 changes are needed for that. 12 If not, what I'm going to suggest is that we reserve any formal actions on this until our 13 14 work session tomorrow. That'll give you 15 additional time to think about any of the 16 parameters, as well -- yeah, when you're done 17 reading your minutes and... DR. ANDERSON: Let's move something. 18 19 DR. ZIEMER: Well, if you want to move 20 something today, you can. 21 We also -- keep in mind -- I would suggest, if 22 you're ready to accept the document, that we do 23 it in a provisional manner, soliciting the 24 input from the Secretary's office as -- and --25 and having a caveat that it's open to

1 additional amendments as the Board sees fit. 2 DR. MELIUS: I would agree with that if -- to 3 use some of my Chicago background -- if you 4 would take -- Paul, if you will take 5 responsibility for delivering Wanda's vote tomorrow. I don't want her being able to 6 7 change her mind overnight. 8 DR. ZIEMER: Well -- well, let me say this. 9 -- if you want to take action on the document 10 itself today, we can do that. I would suggest, 11 nonetheless, that we wait till tomorrow on --12 on identifying the sites that we may wish to 13 have work done. 14 DR. WADE: I think it's better 15 (unintelligible). 16 DR. ZIEMER: If so, I'll entertain a motion for 17 provisional acceptance of the document. 18 that I mean we would consider it our working 19 document for the time being, open to 20 modifications as -- or at least some input from 21 the Secretary's office, as well as --22 DR. DEHART: Second -- I'll second Wanda's 23 motion. 24 DR. ZIEMER: -- as well as possible additions 25 on this issue of using other site data.

1 DR. WADE: Yeah, I think the -- the thought of 2 getting the working group together to -- to 3 complete that issue I think is a good thought 4 and I would hate to see that lost. 5 DR. ZIEMER: So that what it would do would be 6 basically to accept what we have here as a 7 starting point and keep the door open for 8 changes. If the Board is comfortable with 9 that, that would be a motion for something 10 similar. 11 MS. MUNN: Yes. 12 DR. ZIEMER: I hate to put words in other 13 people's mouth, it's a very unsanitary way of 14 speaking. 15 MS. MUNN: Although I'm not the chair of the 16 working group, who perhaps should move --17 DR. MELIUS: I will defer -- I will gladly 18 defer to you making the motion. 19 MS. MUNN: Would you please record that? 20 a first. 21 I move that this body accept the document that 22 is before us as a provisional document, with 23 the understanding that the input of the 24 Secretary will be used to help expand the 25 document and complete it.

1	DR. ZIEMER: I'll interpret that as including
2	the possible expansion for other related
3	topics, as were if that's agreeable.
4	MS. MUNN: Yes.
5	DR. ZIEMER: And seconded by Roy DeHart. Now,
6	discussion?
7	(No responses)
8	Are you ready to vote? You're longing to vote.
9	MS. MUNN: Please.
10	DR. ZIEMER: Please vote. All in favor, aye?
11	(Affirmative responses)
12	Any opposed?
13	(No responses)
14	Any abstentions?
15	(No responses)
16	Motion carries.
17	DR. MELIUS: (Off microphone) Could we now read
18	back (unintelligible).
19	DR. ZIEMER: I believe now the group is
20	getting giddy. I believe that completes our
21	formal action for today. We are going to
22	return at 7:00 p.m. for a public comment
23	period. Let me ask Lew Wade if he has any
24	additional comments or instructions for us
25	before we depart. No?

Okay, we will recess until 7:00 p.m. Thank you very much.

(Whereupon, a recess was taken from 5:00 p.m. to 7:00 p.m.)

PUBLIC COMMENT

DR. ZIEMER: Good evening, everyone. We have some folks still coming in, but I think we're going to go ahead and start, try to keep on schedule, if we can.

My name is Paul Ziemer -- oh, there are some more chairs, though you may want to squeeze together, at least let people know if there's a chair near you. If you can squeeze together a little bit, too, that can help. We'll try to get some more folks in.

I serve as Chair of this committee. I want to just take a minute or two and tell you what the committee does and what we don't do. It's actually an Advisory Board, and we advise the Secretary of Health and Human Services on the dose reconstruction program that's being carried out by NIOSH, and many of you are quite familiar with that program and are even claimants in that program.

This Board is not the board -- we do not do the

dose reconstructions, and we are not an appeals board for people who have claims that have been denied. Rather we are simply an advisory board to the Secretary of Health and Human Services, looking over the agency's shoulder. We are not part of the federal agency. We are all independent folks. I'm -- I'm a retired professor -- and, incidentally, a former Oak Ridger -- and we are looking over the agency's shoulder to determine the extent to which the program is being conducted in the proper fashion.

So we review some of the dose reconstructions in an audit fashion, as it were. We review site profiles and that sort of thing. And in the course of our deliberations where we come together periodically and hear from the agency and from others how the program is going, we always have an open meeting, and through the open meetings we also gain information about how the program is working -- or, for some people, not working, as it may be.

But I do want you to know that if you have individual concerns about your own claim or a claim of a relative, there are NIOSH staff

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people here to assist you with that. So they do want to take care of the individual claims. And many of you will, I'm sure, tonight tell us about your experiences, and from that we learn certain things about what the problems are with the program, what's not working, what is working and so on. So that's really what we're doing here tonight.

Now I have a couple of lists. I have the starting list here of about 15 people who've asked to speak, and there's another page like this, so I'm guessing there's probably about 30 people who have requested to speak. recognize we have scheduled about an hour and a half of time, so if you are one who are -that's speaking to the group tonight, you can do the math there. Let's see, we've got maybe 30 people and 90 minutes, so it doesn't give -if you're taking ten or 15 minutes, you really are going to be depriving someone else of time. So we ask you, to the extent you feel comfortable, try to be fairly concise in what you do.

Now when -- I'm not going to ring any bells or make you sit down, but we do request that you

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try to be as concise as you can and still convey what you wish to the Board. Okay? And the Board members are here. I'm not going to introduce them individually. Their name tags are there. They come from all parts of the country. They represent worker groups, technical groups and so on. We have a broad representation of folks on the Board. them are retired, some still working, so it's a cross-section of people. We -- different ones have expertise in different things. All of us are -- we're like anybody else, you know. We're dumb in certain things and we think we're smart in others, but that's how it is. Okay. I'm just going to go down the list in order and as you -- if you're comfortable, you can stand here, there's another mike here, wherever you're comfortable in addressing the group will be fine. We do need -- when you come up, need to have you repeat your name for our court reporter -- 'cause we do keep a transcript of everything that is said, and that transcript does go on our web site -- so that we capture the information that's presented tonight.

1 So we'll begin with B. A. Austin. B. A. 2 Austin. 3 (No responses) 4 Some folks may have signed up earlier and 5 didn't arrive yet. If B. A. Austin's not here 6 yet, let's go to Gail Elkin. 7 (No responses) 8 Okay. Well, I'll come back and check on these 9 later. Jack Wolum -- Woolum, W-o-o-l-u-m --10 Jack? 11 MR. WOOLUM: I've already got my questions 12 answered. 13 DR. ZIEMER: You got your questions answered, 14 okay, Jack. How about Jim -- I believe it may 15 be Phelps -- Jim, thank you. 16 MR. PHELPS: Hi, my name is Jim Phelps, for the 17 court reporter. I'm here to speak about my 18 father's experience at the Y-12 nuclear weapons 19 plant over the years. I understand that you 20 all are trying to do a dose reconstruction at a 21 plant that had probably weak records on 22 dosimetry. It also is a complicated site where 23 chemicals interfere with the free oxygen 24 radical damage of radiation. Oak Ridge is more

characterized as a chemical damage site than a

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radiation damage site, and that's one of the problems.

You say you want to use best available science. Best available science in Oak Ridge says that you look at the radiation, you look at the mechanism for radiation. And at Oak Ridge -at ORNL when I worked there, we looked at some of those mechanisms way back in the '80s. what we decided is when radiation passes through a medium you get free oxygen radicals. These upset cells do damage to mitochondrias of cells, and these react and make something called superoxide dimutase enzymes, which go and repair that damage. If that damage gets very high, you start noticing things like mycoplasmas becoming active in the persons and you have to sometimes give them antibiotics to have them recover from high dose radiation. What happens next is you get competition for the trace metals in the cell that disrupts another enzyme called 2-5A RNase L, and the metal that gets in competition is manganese. And what happens is a mutation of that enzyme from a normal 83 kilodalton weight to a 37 kilodalton weight. When that mutation occurs,

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the cells can no longer defend themselves from pathogens, be that pathogen mycoplasmas or the complement of viruses that you'd normally see in cancer tumors. When you lose that mechanism, cancer viruses and bacterias can take over control for the cytokine mechanisms in the cells. Now that's the simple approach to radiation damage.

We didn't stop there at ORNL back in the '80s. We went and added the chemical damage vector component, and one of the things we noticed was that hydrogen fluoride easily enters the body in places like Y-12 in their green salt operation and places like the hydrolysis of UF-6 released from K-25 makes HF. As people are exposed to that, that's a fairly cumulative poison that deposits itself in the tissues and bones and gets progressively worse with age. And what happens is whenever you encounter aluminum in the environment, it will spontaneously combine in the body to form ALF3. ALF3 mimics a hormone called the thyroid stimulation hormone, and when that happens you no longer have the night and day variation that you get that allows your body to rest at night

and do cellular repair. And what it seems to do is deplete an enzyme called glutathione. When that glutathione enzyme becomes depleted in persons exposed to those kinds of chemical toxins, you no longer remove the mercury from your body, and several other metals. And as mercury builds and it gets involved with the mtDNA and causes respiration dysfunction. How it makes ATP, it generates free oxygen radicals that does the same sort of damage as radiation pathways going through the cell.

So what we identified was the chemical damage vector adds directly to the radiation damage vector. It's probably the bigger component at Y-12 from many operations and at K-25 from many operations. So I attended one of your last meetings over in Knoxville and the word was best available science. I think that we found a better available science back in the '80s at ORNL that speaks to this mechanism that I just pointed out to you. And I think, in terms of doing dose reconstructions, that most of these plants need to have the chemical vector put in there, more so than probably the radiation vector in many of these cases. So when you

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just look at radiation, you're doing many of these people a disservice because most of them -- some of these chemicals like PCBs, uranium machinists at Y-12 were, in the early days, exposed to PCBs. PCBs is one of the chemicals that shuts down glutathione. When you lose that -- they also had a lot of mercury vapors at Y-12 -- you get mercury buildup, you get this reactive oxygen species damage generated in your cell mitochondria and it adds to the radiation component, so -- and it's usually much bigger, and it's a more cumulative process 'cause PCBs are fairly retained, hydrogen fluoride's fairly highly retained, more so than some of the internalized uranium and various other internalized isotopes and external radiation kind of problems.

So that's what I wanted to point out. I would like to ask that you somehow figure how to include that sort of criteria in doing these folks' dose assessments. It needs to go beyond just radiation because, you know, in the '80s we figured out this better model, this better modeling system that everybody that was in the national security group that I worked with that

1 was doing some -- trying to figure out what 2 went wrong in Oak Ridge -- and agreed with it. 3 It was standard available science in the 1980s. 4 It has only been proven harder and firmer and 5 difficult to ignore in this day and age. 6 So anyway, thanks for your time. 7 DR. ZIEMER: Thank you very much, and certainly 8 that is a challenging issue to consider. 9 would point out to the Board that I have 10 received from Mr. Phelps a detailed copy of a 11 letter which I think delineates pretty much 12 what he has provided, and I will make sure that 13 the Board members get a copy of this, as well, 14 Mr. Phelps. 15 Oh, have we distributed that already? 16 we'll make sure the Board members get copies of 17 that, as well. Thank you very much. 18 Then Otis Lee. Otis Lee? 19 MR. LEE: I pass. 20 DR. ZIEMER: Pass? Okay. T. L. Disman. 21 MR. DISHMAN: Hi, I'm T. L. Dishman --22 DR. ZIEMER: Dishman, okay. I said it wrong. 23 MR. DISHMAN: -- retired from Y-12 after 37 and 24 a half years. I'm real disturbed at this dose 25 reconstruction because after 37 and a half

1 years and wearing a dose meter for maybe ten 2 days during 37 years, I'm one that would know 3 you couldn't reconstruct my record. And I 4 think everyone in this room knows you can't 5 reconstruct records that don't exist. 6 we've suffered with this malfeasance for all 7 these years, and we're going to continue --8 maybe our -- maybe our speaker that just 9 finished has some good suggestions for us. 10 Maybe we'll just check your health instead of 11 your record 'cause you've got your health 12 problems, but your records don't exist. So we 13 need to not let everybody die before we say 14 gosh, that's what we should have done. It's --15 there -- there's a sin going on here, and it's 16 a sin against these people, and it needs to be 17 corrected and it needs to be corrected very 18 Thank you. soon. 19 Thank you very much. DR. ZIEMER: 20 UNIDENTIFIED: Mr. Zimmer (sic), can they stand 21 over here? 22 DR. ZIEMER: Yes --23 UNIDENTIFIED: (Off microphone) 24 (Unintelligible) stand over here --

DR. ZIEMER: -- maybe -- maybe, Dr. Anderson,

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1	if you could help move the podium and then they
2	
3	UNIDENTIFIED: (Off microphone)
4	(Unintelligible) see the (unintelligible)
5	DR. ZIEMER: Yeah, sure, we can do that. We'd
6	sort of like to see their faces, too, but we're
7	facing each other. How about sideways, will
8	that help to sort of Thank you.
9	Okay, next we have Cleveland Drummand.
10	UNIDENTIFIED: We want to see everybody's face.
11	Would you move the podium completely away?
12	DR. ZIEMER: Oh, sure, yes.
13	UNIDENTIFIED: We're looking at you.
14	DR. ZIEMER: Yes, I gotcha. Okay.
15	(Pause)
16	Cleveland Drummand?
17	UNIDENTIFIED: (Off microphone)
18	(Unintelligible)
19	DR. ZIEMER: How about Lester is it Branham,
20	Lester Branham?
21	UNIDENTIFIED: (Off microphone)
22	(Unintelligible)
23	DR. ZIEMER: Oh, okay. Ray Beatty? Ray
24	Beatty.
25	MR. BEATTY: Good evening. My name is Ray

Beatty. I'm a 14-year employee at the Fernald site in Fernald, Ohio. I've come here tonight to express some opinion as to the Board membership. I was involved in the conference call that you all had about a month ago, and I heard of the new Board members coming on board, so I intend to address that tonight, and I think you saw today that there is an SEC petition, number 46, for the Fernald site, and this information would be very applicable to that application.

It's in reference to Dr. James Lockey who's been appointed to serve on the Advisory Board, beginning with the next meeting. Having attended many of these meetings, I am pleased to see him here. Dr. Lockey's had extensive involvement at Fernald. He serves as the government's appointee to review causation related to state workers compensation claims as part of the Fernald Settlement Trust arising out of the case David Day versus (unintelligible) Industries of Ohio.

There's a three-physician panel for Fernald

workers which reviews workers' cases to

determine whether DOE or their contractor will

1 contest the claim. Dr. Lockey was also 2 appointed by DOE and its contractor, Lockheed-3 Martin, to review 55 six worker claims in Oak 4 Ridge as part of the Lockey/Byrd/Freeman panel. 5 I am requesting that Dr. Lockey's conflict of interest forms, his waiver letter and the 6 7 relevant elements of his personal financial 8 disclosure form that deal with his expert 9 witness and other government DOE contractor 10 consulting work be made public. We have no 11 interest in his personal financial matters, 12 only matters affecting public service. 13 However, he should make public a list of all 14 workers compensation or court cases -- pardon me, tort cases, claims where he has been served 15 16 as an expert witness, whether testifying or 17 not, and involving DOE or its contractors. 18 Dr. Lockey's reportedly had consulting 19 arrangements in defense of litigation claims on 20 behalf of various industries. This, too, 21 should be disclosed as it may impact his public 22 service. 23 In addition, I would hope that Dr. Lockey will 24 have no part in any deliberations, votes, 25 reviews involving Fernald SECs, site profiles,

1 dose reconstructions or technical documents. 2 While Dr. Lockey may be a fine person and 3 someone supported, the integrity of the program 4 for Fernald workers will be impaired if he's 5 allowed to take part in any matter impacting Fernald workers. NIOSH has no shortage of 6 7 staff that worked at Fernald who can contribute 8 to their expertise. Information posted on the 9 NIOSH web site for conflict of interest is far 10 too limited to describe the full range of 11 conflicts that should be made available. 12 would hope that if Dr. Lockey's permission is 13 needed to disclose some of this information, 14 that he would provide the necessary permission. 15 Thank you. 16 DR. ZIEMER: Thank you very much, and just as a 17 general statement I'll point out that we do 18 require all of the Board members to have a 19 conflict of interest information on the web 20 site. I believe -- the new members are 21 probably not there yet, but --22 MR. BEATTY: They're -- they're not there, I've 23 looked. 24 DR. ZIEMER: -- they're not there yet 'cause 25 they don't begin their term --

1	MR. BEATTY: And I have read
2	DR. ZIEMER: till next month.
3	MR. BEATTY: each member's conflict of
4	interest and saw where a couple had a conflict
5	of interest, but this one is much more
6	extensive that I just alluded to.
7	DR. ZIEMER: Yes. It will certainly be on the
8	web site as soon as he begins his term, and we
9	do have a Board rule on conflict of interest
10	issues where we're not allowed to vote on sites
11	where we have those conflicts, so
12	MR. BEATTY: Okay, I we (unintelligible)
13	that.
14	DR. ZIEMER: we appreciate your heads-up on
15	that.
16	MR. BEATTY: Thank you very much. Thank you.
17	DR. ZIEMER: Thank you. I'm having a little
18	trouble reading this one. It may be Johanna
19	Goodman, or it Okay, good.
20	MS. GOODMAN: My name is Johnnie Sue Goodman.
21	DR. ZIEMER: Oh, Johnnie? Okay
22	MS. GOODMAN: Uh-huh.
23	DR. ZIEMER: Johnnie Sue, thank you.
24	MS. GOODMAN: And I am here on behalf of my
25	late husband. In 1980, January of 1980 he took

sick all of a sudden. Now he'd been feeling bad for several months, but we didn't know what was wrong with him and he wasn't one to run to a doctor every time he got -- got a little bit under the weather. Well, in January of 1980 he took a hurting right in here, right in his chest. He thought maybe he had the flu or something.

He went to the doctor over at Oak Ridge at work

that day. He said oh, take a couple of Bufferin and you'll be okay in a few days.

Well, he continued to get worse for about three or four days. I took him to an old doctor that -- up at Concord, Dr. Malcolm -- whatever his name was, but anyway, that slipped my mind just then. But he said what's wrong with you? And he said I -- I can't tell you, let my wife tell you. So I went through the ordeal of it and he said well, let's step out in the hall a few minutes and I will send the nurse in to do some blood work on him.

So he took me out in the hall and he said

Honey, I hate to tell you, but your husband is

almost dead. I said well, how -- how come? He
said he doesn't have any blood. He said he's

1 got leukemia in the worst way. And I said well 2 -- well, how do you know? He said well, I've 3 seen enough of it -- it's work-related. 4 I had no idea what that meant. My husband had 5 worked over there over 20 years. He had never 6 told me anything that he ever done except he 7 was an assistant general foreman in the machine 8 shop. I had no idea what he did. 9 So it -- I was in shock. Well, we got him in 10 the hospital. In 60 days he lost 60 pounds. 11 They gave him 150 pints of blood. It didn't do 12 any good. They tried every kind of chemo on him they could. They -- I thought -- was 13 14 trying to kill him. He was so pitiful, he 15 couldn't eat. He was in misery. And on the --16 July the 12th he passed away -- well, we buried 17 him on the 12th. He -- he died on the 9th of 18 July. He suffered death many, many times. 19 I have never got any compensation, but men like 20 that -- not only my husband, but others -- they 21 need to be repaid some way or other, and I can 22 -- I need it, and that's all I have to say. 23 DR. ZIEMER: Thank you. And thank you, 24 Johnnie, for sharing that with us. I know 25 that's difficult.

1 Helen -- let's see, I'm having a little trouble 2 reading the last name -- it looks like G-a-l-h-3 s-o-n, Galhson? Probably don't have the last 4 name correct. 5 Oh, Helen G -- maybe it's an Alban or Allison -6 - any Helens? Any Helens? 7 **DR. ANDERSON:** What's the address? 8 DR. ZIEMER: The address is -- okay, we're 9 getting it closer. I'm sorry, Helen, I'm just 10 having a little trouble reading this. UNIDENTIFIED: Well, I didn't come here to 11 12 speak, but I'm just like her. My husband got 13 sick and didn't know what was wrong with him 14 and he only lived three months and they found 15 out he had acute leukemia. And as you say, 16 after 37 years, I think we all deserve a little 17 bit, and we're not getting too far. 18 DR. ZIEMER: Okay, thank you. 19 UNIDENTIFIED: Thank you. 20 DR. ZIEMER: Then I have Franklin Tucker. 21 MR. TUCKER: Gentlemen, my name's Franklin 22 Tucker. I worked at Y-12 approximately 12 23 years before they forced me to retire back in 24 about 2001. I was a chemical operator. Now 25 you're talking about doing a reconstruction

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claim -- or reconstruction and stuff on doses and stuff like that. You're going to run into some serious problems. Like other people I've heard here talk about records and stuff, what don't exist or what they have made up to fill in gaps. I see no way of ye'uns (sic) compensating for that. The whole 12 years I was there working in enriched uranium and also in special materials organization where the chemical exposure got me, there was never no study done. There was nothing done. Now I have been fighting this approximately three and a half years now. I went all the way to President Bush because I worked on weapons systems like the 88, the 87 and stuff like that. And when you work in that stuff and you start bleeding out the nose, and you're so sick that for the next couple of days that you can do nothing, something is seriously wrong. I went to the Department of Energy, voiced my complaint. All they said was hey, you're a whistle-blower. When I finally did come down sick and they told me I had to go out on longterm disability, I was told by the people at Vanderbilt and stuff like that I have

1 what's called a chronic wasting of the brain. 2 My brain's dying. 3 These people -- I mean -- and since I've been 4 sick -- and this is the sad part of it, the 5 people I have run into that are so sick and 6 have to fight the system to get anything. I 7 have not been -- I mean I have been treated 8 very bad, like I said, since I become sick and 9 stuff like that. 10 Ladies and gentlemen, the thing I recommend, 11 one thing is please contact the President of 12 the United States. You can get his phone number. Call him. Tell him, say you know, I 13 14 worked on such-and-such weapons system, and 15 that will get your atten-- his attention right 16 there, and tell him what happened. And then 17 request -- say where is a reconstruction on our 18 dosage and limits and stuff of chemical 19 operators and machinists and stuff, the two 20 people that would have been exposed the most, 21 'cause there's never been a study done. 22 checked. 23 That's all I have to say. Thank you. 24 DR. ZIEMER: Yeah, thank you very much. Colleen Schotz?

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1 MS. SCHOTZ: I'll pass. 2 DR. ZIEMER: Okay, thank you. Let me jump back 3 to the beginning of the list again. B. A. 4 Austin, did he come in -- or she? 5 (No responses) 6 How about Gail Elkin? I know in some cases the 7 request for public comment sheet was 8 inadvertently signed by people who thought they 9 were simply registering, so some of those may 10 have been in that category. 11 I now -- I now need the second sheet and we'll -- I think the second sheet were -- were all 12 13 folks who came here this evening, so -- yes, 14 sir? 15 UNIDENTIFIED: I'd like to find out if you can 16 tell us who these people are at the table. 17 DR. ZIEMER: Yes, we can -- we can go around 18 the table. I indicated that these are the 19 members of the Advisory Board. Dr. Henry 20 Anderson, who's from Wisconsin -- maybe I 21 should have each one tell who they are and where they work. Why don't -- Mike, why don't 22 23 you go ahead and -- this -- this is the 24 Advisory Board on --25 (Whereupon, several members of the public began

1	speaking at once, without benefit of
2	microphone, rendering none of the comments
3	distinguishable enough for transcription.)
4	DR. ZIEMER: Sure. Sure, sure.
5	MR. GIBSON: Hello, my name is Michael Gibson.
6	I worked at the Mound facility,
7	(unintelligible) for Ohio for 23 years. I was
8	an electrician by trade. I was also a union
9	president for probably ten years. I was
10	appointed to the Board in August of 2002.
11	DR. ZIEMER: Thank you.
12	DR. DEHART: I'm Roy DeHart. I'm currently at
13	the Medical School of Vanderbilt. I grew up in
14	Oak Ridge. I worked at X-10 and Y-12.
15	MR. ESPINOSA: Richard Espinosa, currently
16	employed at Los Alamos, New Mexico. I work as
17	a sheet metal worker and chief steward for Los
18	Alamos.
19	MR. GRIFFON: I'm Mark Griffon, a health
20	physicist, and I'm currently involved in a lot
21	of medical surveillance programs around the DOE
22	complex.
23	MR. PRESLEY: Most of y'all know me. I'm Bob
24	Presley from Oak Ridge. I worked at Y-12 37
25	and a half years. I'm back out there.

1 DR. ZIEMER: Well, again -- I'll go ahead and 2 reintroduce myself, some weren't in here. My name is Paul Ziemer. I'm a retired professor 3 4 from Purdue University. My area is health 5 physics, and I got my start in my career at X-10 and a little bit at Y-12, as well. 6 7 DR. WADE: My name is Lewis Wade and I work for 8 NIOSH, the Centers for Disease Control. 9 the Designated Federal Official on the Board 10 representing the Secretary of HHS. 11 DR. MELIUS: I'm Jim Melius. I'm a physician. 12 I work for the laborer's union. I'm Wanda Munn. I'm a retired 13 MS. MUNN: 14 nuclear engineer from Hanford. 15 DR. ROESSLER: Genevieve Roessler, I'm retired 16 faculty from the University of Florida. 17 field is health physics. I'm now living in 18 Minnesota. 19 DR. ZIEMER: Okay, thank you. Let's continue. 20 Then we have Dorothy Thompson. 21 MS. THOMPSON: Well, I'm another Y-12 widow, 22 and I guess it's very clear to most of us in 23 this room that Y-12 is as dangerous as -- was 24 as dangerous, is as dangerous, I don't know 25 what the situation is -- as any other place.

1 My husband died after a five-week illness of 2 cancer in 19-- in 2001. The primary site was 3 never really discovered, as apparently that --4 it had gone everywhere. It was an unusual 5 variant of a patocellular cancer. 6 I applied and was just recently turned down by NIOSH. They gave me a 45 percent. 7 I don't 8 think it's accurate. I don't think it's fair. 9 And I don't think it -- it's fancy algorithms 10 that fancy mathematicians have done, but I 11 don't think it tells the story. 12 My husband was employed in 1961 as a special 13 project engineer. He was in the 18-day 14 turnaround program at Y-12 where they would get 15 orders from Los Alamos or someplace to build 16 parts. He would stay out there on weekends. 17 The plant manager, Jack Case, saw that he was a 18 bright young engineer. He would stay out there 19 on weekends with a health physicist outside the 20 door and make parts, carry parts, examine 21 parts, inspect parts, and was promised that all 22 the rest -- everything else that he did not 23 work with would be buried. And he would say 24 Honey, you don't know what I've done today. 25 You don't -- don't let the kids touch my shoes,

don't let -- you just don't want to know what I've done. But he said they put a lead apron on me most of the time, and Case thinks I'm really a nice young man and I've got places to go. And he lost his life for Y-12.

Now interestingly enough, he was in this 18-day turnaround program from 1961 to 1967. Our first child, born in 1964, was normal. Our second child, born in 1968, had a devastating, etiology unknown, birth defect. Our third child, born in 1970, had a devastating birth defect.

NIOSH says it's -- oh, we received counseling from the Mayo Clinic, from the University of Tennessee, from Vanderbilt, and they all said do not have any more children; we fear that your husband's sperm has been radiated.

He was -- Bill was also in charge of the mercury cleanup at Y-12 for a long time.

Gentlemen, it's not fair. The -- oh, so I went to a lawyer, and the lawyer said you know, if you can prove he had any other kind of cancers, because of this 45 percent, you're probably okay. Well, Bill did have other cancers. But guess what? It took from 19-- 2001 to now,

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2006, to get my rejection from NIOSH. And guess how long doctors keep their records? They've all been shredded. They've all been shredded. You can call the family clinic, they've all been shredded.

Dr. -- the doctor on -- Dr. Sharp, who did all of his skin cancers, there are no more records. All I have -- I have a bill. I don't know what -- I don't know where to go. I don't know what to do. I was told by the lawyer also that if I could have proved he had worked at K-25 for as long as six weeks, there would be no problem, I would get the money. And it's really not about the money. I lost my husband at 63, and he was a good man. He worked with Bob. And -- and he -- the grandchildren are without a father, my brain-damaged daughter is without a father. It's not fair, guys. It's -- the NIOSH simply does not adequately reflect individual spikes, individual incidents, and I'm told that I can't question NIOSH, that the only way I would get the money is if I could prove he had more cancers, which is my word against your word, or if he'd worked at Y-12 -- I mean if he'd worked at K-25.

1	So I think somebody needs to relook at the
2	process because there's no way that dose
3	reconstruction adequately accounts for where he
4	was from 1961 to 1967. Thank you.
5	DR. ZIEMER: Thank you, Dorothy. Next I have
6	Thomas Duncan.
7	MR. DUNCAN: My name's Thomas Duncan. I met
8	y'all a few months back.
9	DR. ZIEMER: Yes.
10	MR. DUNCAN: And the only way I got to go to
11	your meeting was big issue, they you have
12	to have authorization to get off work
13	DR. ZIEMER: (Unintelligible), uh-huh.
14	MR. DUNCAN: and I tried to get
15	authorization this time, they denied it. And
16	that's the reason I haven't attended any of the
17	day meetings.
18	DR. ZIEMER: Yeah.
19	MR. DUNCAN: I got five different kind of
20	cancers. I worked in 18-day turnaround for 12
21	years. It's deadly. I got in contact with
22	NIOSH my wife called me at work Friday and I
23	called them Friday. I got the letter they sent
24	me Saturday and read it, and of course getting
25	authorization, they told me last time through

labor relations and my division -- department head, I have to give them opportune time and to get that information for the last meeting took me about three weeks for them to give me that denial for the last one. And if you'd like more people to attend your meetings, you know, I consider it company business, you know, if it's -- you're talking about Y-12. Maybe get in contact with Y-12 and -- and let the people off, you know. If -- if I took anything besides vacation, they said it'd be disciplinary action up to termination, and that's from labor relations.

And for NIOSH, I've contacted them two or three times and they said well, we'll give you a call back, never hear nothing from them. And I asked for some records where I attended the last meeting, you know, I signed in on the books, and they said yeah, we'll have them sent to you, and that's been three months ago, still waiting on them to send them to me.

I got a letter today from NIOSH on my update and I've had some other body parts removed just a month or so ago, and I sent them in to my representation and I think -- I think I got

1 five different kind of cancers, and that make 2 the sixth one, and it's not showing up on my 3 records through NIOSH, and I don't know how 4 long it takes for them to -- you know, I been 5 fighting cancer for oh, a little better than a year, and they say oh, you've got the best 6 7 kind. You know, it's 80 percent curable. 8 Well, I ain't reached 80 percent yet. 9 about all I can say. 10 DR. ZIEMER: I wonder if some of the NIOSH 11 staff can make sure we get the information from 12 Mr. Duncan for the records and whatever was There's some staffers here that can 13 requested. 14 help you yet tonight, I think, and try to get 15 that for you. 16 Actually with the number of people who declined 17 to -- to speak, we actually have finished the 18 list here, so I'm going to ask -- there may be 19 others who didn't sign but that do wish to 20 speak, and I'll give you the opportunity. 21 Yes, sir, please approach the mike and give us 22 your name. 23 MR. ROYSTER: My name's Paul Royster. My dad -24 - some of y'all might know my dad. His name 25 was Billy Royster -- George William Royster.

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He died in 1968. Our claim was denied for -said it had 48 percent. Some of the things he was involved in was he was there when the atomic bomb ground testing in 1957. He was involved in some experiments where he drank radiated milk from cows. Frank Munger* did a story about this a while back. He had plutonium spilled on him, that's documented. It spilled on his hands. That's documented. Some of his documentation was just handwritten, you know. He worked at UT Farm and he also worked at Y-12. He was a health We've been involved in the same -physicist. same fight with these other people. He also had two years of documentation that couldn't be found from UT Farm. But anyway, he died in 1968 when I was nine years old. -- my mom got \$25,000 life insurance to raise five kids, so I'm really here on behalf of my She hadn't really been involved in this that much. My brother mainly has been involved with this, but the case number is 1407 and the dose reconstruction, 48 percent up to 50, I don't see how you can vary two percent. There should be a leniency there, to me. This lady

said 45 percent. It seems like there should be like a five or ten percent one way or the other, the way I see it.

But anyway, I haven't really been involved in this that much, like I said. My brother has, but like I said, the experiments with the radiated milk from the cows and there were five other people involved in that. I don't know how long that went on, but -- then when the atomic bomb was dropped and they just said put some glasses on and turn your head, you know. And he died of a brain tumor in 1968, but that's just -- I just wanted to share that with you.

DR. ZIEMER: Thank you very much. Yes, please, ma'am, if you wish to...

MS. ROBERTSON: My name is Florene Robertson, and I'm here on behalf of my husband, who passed away almost 11 years ago. He passed away with cancer of the colon and of the liver. Now he worked at X-10. Some of the people here may think X-10 is clean, but there are contaminated areas at X-10. And there was a night that he worked overtime. In fact, it was on a Saturday. He went in and worked on

1 Saturday, and he called me and says I'll be a 2 little late, I'm working over. Okay, he did 3 not come home until the next morning, Sunday 4 morning. And he had been there all night long 5 doing a what they call wash-down. They had to 6 wash him down, so that man was contaminated. 7 All right. The fact that I want to tell you 8 that that area's also contaminated is that they 9 -- he was on loan at different times. He would 10 come home and I'd say well, what'd you do 11 today? He'd say oh, I worked at Y-12. So I 12 don't have any way of proving that he worked at 13 Y-12, but he did work at Y-12 and he also 14 worked at X-10, but he was classified as a pipe-fitter. And in the area that he worked in 15 16 I'm sure was pretty safe, but he did contact 17 radiation while he was there. Thank you. DR. ZIEMER: Thank you. Yes, ma'am, uh-huh. 18 19 MS. BURGESS: Good evening, thank you all for 20 being here. I'm here --21 DR. ZIEMER: Give us your name, too, for --22 MS. BURGESS: Oh, I'm sorry --23 DR. ZIEMER: -- the record. 24 MS. BURGESS: -- my name is Gail Burgess --25 Gail Mynant* Burgess. I'm here on behalf of my

1 father, who I lost 30 years ago to eye cancer. 2 And I have no guarantee that the records that 3 you guys are looking at are true, are factual. 4 I can't get in to see them. They've all been 5 shredded. They're missing. His medical records are missing, even at M. D. Anderson 6 7 where he went twice. And just as a personal 8 note, I worked in the field at X-10 for Bechtel 9 National, and we took samples. I didn't get a 10 dosimeter for two years. And then when I left 11 and went back to the tower in Oak Ridge, they 12 didn't do a full body count, so they don't even 13 know how much radiation I got. 14 Tell me now -- you've got me working out there 15 12 years ago, and then my father working out 16 there 30-some-odd years, I don't understand how 17 you people are going to put it all together. 18 And how can you let this woman and this man 19 have -- be so close and not be paid? I don't 20 understand this. There's got to be another way 21 to do it. 22 DR. ZIEMER: Thank you, Gail, and yes, ma'am, 23 go ahead. 24 MS. SLACKEY: My name is Sharon 25 (unintelligible) Slackey. First of all, I want

to say that every one of us wouldn't be here if we didn't have a story. All of them are alike. I want to know what are you going to do about them? Do you have any power at all to do anything?

I had a father who is dead. He died of cancerous brain tumor. He worked at Y-12. They said they weren't responsible in the dose reconstruction. They had many years where he didn't even have a dosimeter, and yet he crawled in and out of the pressure vessels at 9-212, and I do know what the pressure vessels at at 9-212 were used for because I had a son that was there that was regularly exposed. Had he not died in a single-car accident, he would have died of cancer also.

I have been retired from Y-12 now for about three years. I have also gone through cancer and I'll tell you right now they'll probably turn me down. They turned us down on our father, and our father had a cancerous brain tumor while he was working there. They did not diagnose it until six months before he died. They didn't know to look for it. But he got it at Y-12. We all know that.

1 My husband was a subcontractor out there at Y-2 12 and did -- was doing a job in the east 3 ponds. They let the -- turned the water loose 4 on him and it was hotter than a depot stove. 5 He was irradiated. He's a three-time cancer survivor. There's no medical records to back 6 7 it up. That all happened a few years ago. 8 If you guys can do something about these 9 stories, it'll make these meetings worthwhile. 10 If you can't, I don't see why you're having 11 them. 12 DR. ZIEMER: Yeah. Thank you. I might -- I might comment that with -- there is a sense in 13 14 which what we're able to do is somewhat 15 limited, but we are trying to do what we're 16 able to do, within what the law allows us to 17 do, to address some of these problems. 18 hopeful that in cases such as yours where there 19 are missing records that there may be alternate 20 ways to establish the situation. We will not 21 always be successful --22 UNIDENTIFIED: (Off microphone) 23 (Unintelligible) disease, but you know why they 24 couldn't pay for that?

Yeah.

DR. ZIEMER:

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And

1 **UNIDENTIFIED:** He died in 1981 and he had to 2 have had the test that was given in 1993. 3 DR. ZIEMER: Yeah. Thank you. 4 UNIDENTIFIED: You're damned if you do and 5 damned if you don't. 6 DR. ZIEMER: Okay, ma'am -- yes, please. 7 MS. MILLER: Hello, my name is Kathy Miller, 8 and I'm here -- I'm also a retired nuclear 9 worker, but I'm here on behalf of my father's 10 claim. And I have -- this is not unfamiliar to 11 me, having worked in a government facility. 12 I'm caught in a bureaucratic loophole and no 13 one will take responsibility between NIOSH and 14 the Department of Labor, and I thought maybe 15 you all could find the answer to this question. 16 My father went straight from the south Pacific 17 at the end of World War II into work at Y-12, 18 and he -- this is the week of his 25th 19 anniversary of his death. He died -- he was 20 diagnosed at age 54 with multiple myeloma, and 21 he died be-- when -- he died before he was 60, 22 he died when he was 59. So my mother was 23 without his companionship and his help --24 earning help and all those things for all those 25 years, and she filed in November of 2001.

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she lasted as long as she could, but she died about a year ago when she was in her eighties. So this is my question. My father's record, I'm told by NIOSH and by the Department of Labor, has been pulled as part of the special dose cohort or something along that line because he came to work at Y-12 in 1945. the people who are -- who were -- I believe, is this correct -- worked for -- entered work from '43 to '48, those records have been pulled and set aside to be evaluated separately by the Department of Labor. Now we've been waiting since November of 2001 for my father's dose reconstruction and we don't have it, so now at the end of September, this law that he was covered by went into effect, September 25th, and at that moment or shortly thereafter NIOSH pulled these records -- I think there's several hundred of them -- and ceased work on the dose reconstruction and forwarded them to the Department of Labor.

Okay. They have been there at the Department of Labor since that time or around the first of the year -- I know how things -- slowly things move. At the Department of Labor they tell me

1 they're waiting for guidelines to administer 2 this Act, and they -- I've tried to find out 3 who they're waiting on so I could say, you 4 know, could I get in touch with them, is it a 5 Congressman, is it a Senator, is it a committee, is it a staff, what's the situation; 6 7 they don't know. And I've called back to NIOSH 8 and they say no, we're not doing it. So right 9 now, since September, all these several hundred 10 have just been sitting there, nothing's being 11 done on them. And I've called about every two 12 weeks since I found this out, and the 13 Department of Labor has no new information. 14 So I got really frustrated about two weeks ago, 15 went to Senator Lamar Alexander's office and a 16 staff person there has been very kind and been 17 trying to help me, and she's been unable to 18 find out. 19 So my question is, I'm willing to push on 20 somebody to do something if I know who it is 21 and where to go. 22 DR. ZIEMER: Let's see if we can find someone 23 who at least knows where she should be 24 directed, either --25 MS. MILLER: I think there's 800 cases involved

1 in this. 2 DR. ZIEMER: Okay, right -- right here in the 3 back, from NIOSH --4 UNIDENTIFIED: (Off microphone) 5 (Unintelligible) DR. ZIEMER: -- or Department of Labor -- we 6 7 have both NIOSH and Labor here --8 MS. MILLER: Okay, thank --9 DR. ZIEMER: -- and at least someone that is in 10 the position to answer that question. 11 you. 12 Okay, ma'am, please. 13 MS. HOLT: My name is Faye Holt. I lost my 14 husband in 1954 with cancer. And I'm concerned 15 who pays the salary for you guys to be here? 16 DR. ZIEMER: Let me address that --17 MS. HOLT: Would all of you --18 DR. ZIEMER: No -- yes. 19 MS. HOLT: -- like to know? 20 DR. ZIEMER: Yes, you would like to know that. 21 First of all, let me tell you that it does not 22 come out of the pot of money that's used to pay 23 claims. No, some people think that their claim 24 money is being used to pay people such as us. 25 It -- that's a separate pot of money and that -

1	- that money is there, regardless of whether or
2	not, for example, this Board meets.
3	These Board members are on the federal
4	they're considered special federal employees.
5	We get paid the federal consulting rate, which
6	I can tell you is about one-tenth of the
7	commercial rate, and I don't know what I'll
8	tell you what I made this year in serving on
9	this Board. I made \$5,500.
10	MS. HOLT: Well, you have all of you have
11	jobs and you're just
12	DR. ZIEMER: I'm retired.
13	MS. HOLT: volunteering to do this?
14	DR. ZIEMER: I'm retired. I'm retired, so I
15	supplemented my my Social Security by that
16	amount. And
17	MS. HOLT: None of you then are employed by
18	NIOSH or DOL?
19	DR. ZIEMER: No. No.
20	UNIDENTIFIED: (Off microphone)
21	(Unintelligible)
22	DR. ZIEMER: Well, Lew Lew Wade is a
23	Designated Federal Official, which is a federal
24	requirement for this
25	MS. HOLT: You're with NIOSH, correct?

1 DR. WADE: Yes. 2 MS. HOLT: Well, I have a question for you. 3 How can you do dose reconstruction on an 4 employee if you do not know whereabouts, what 5 building -- you know they worked at Y-12, K-25 6 and X-10, but you don't know whereabouts and in 7 what building they worked? Then how can you do 8 dose reconstruction based on coworker 9 comparison? There's no way you can do it if 10 you don't know where the man worked. He could 11 have worked up in a ceiling, he could have 12 worked under the floor. Where are you going to 13 go to get someone to compare with his case, 14 with his --15 DR. ZIEMER: Let me take --16 MS. HOLT: -- (unintelligible) --17 DR. ZIEMER: -- Mr. (sic) Wade off the hook 18 because he is not actually a member of this 19 Board, nor is that his area of expertise. But 20 21 MS. HOLT: Well, is (unintelligible) --DR. ZIEMER: -- in fact --22 23 MS. HOLT: -- can answer that? 24 DR. ZIEMER: -- in fact, the challenge -- the 25 challenge that NIOSH has is to do what you

1 described, and if they are not able to do that, 2 then --3 MS. HOLT: They estimate. 4 DR. ZIEMER: Well, if they --5 MS. HOLT: Yes. DR. ZIEMER: -- if they estim-- if they cannot 6 7 estimate within reasonable scientific bounds, 8 and one of the jobs of this committee is to ask 9 that question, whether they are in fact doing 10 that, and if they're not able to reconstruct 11 doses or make a scientifically-defendable 12 estimate, then they have to --13 MS. HOLT: They assume --14 DR. ZIEMER: -- place -- they have to place the 15 individual in what is called the Special 16 Exposure Cohort. And of course that really is 17 the issue that is being --18 MS. HOLT: But how can they assume --19 DR. ZIEMER: -- being struggled with with Y-12 20 is in fact can -- can what you describe be done 21 22 MS. HOLT: But how --23 DR. ZIEMER: -- yeah. 24 MS. HOLT: -- can they assume that he can be 25 compared to Joe Brown when they don't know

1 where he worked? 2 DR. ZIEMER: That is -- that indeed is the challenge. And if it can't be done --3 4 MS. HOLT: Well, they say they know he had 5 various exposure to radiation, but they don't know where he worked. So now how do they know 6 7 he had various exposure? 8 DR. ZIEMER: Well --9 MS. HOLT: I mean, you know, there needs to be 10 11 DR. ZIEMER: -- we -- we --12 MS. HOLT: -- some answers. 13 DR. ZIEMER: Actually -- and the answer is 14 actually fairly lengthy, but --15 MS. HOLT: Well, I've got all the time you 16 want. 17 DR. ZIEMER: -- but what I was going to suggest 18 -- what I was going to suggest is that we could 19 -- we could take some time and ask, for 20 example, one of the NIOSH people to give a 21 quick overview of that process, if the group 22 would like to do that. We --23 MS. HOLT: Well, all of the people that are 24 here tonight, we didn't come up here for a 25 picnic or a piece of coffeecake and --

1 DR. ZIEMER: Understood. MS. HOLT: -- a cup of coffee. We came up here 2 3 to get some things done. 4 DR. ZIEMER: Right. 5 MS. HOLT: Well, evidently you all are not 6 doing any more than what the letters say that 7 we get, so why come to a meeting? Right? 8 DR. ZIEMER: I'm wondering if --9 MS. HOLT: Is everyone in agreement? 10 DR. ZIEMER: I'm wondering if Jim Neton --11 MS. HOLT: What we all need to do is join --12 DR. ZIEMER: I'm going to -- I'm going to put 13 Mr. (sic) Neton on the spot. Dr. Neton is the 14 -- sort of the chief guy for NIOSH for dose 15 reconstructions, and he will describe briefly -16 17 MS. HOLT: He's doing a terrible job. Well --18 DR. ZIEMER: 19 MS. HOLT: No one is doing anything. 20 DR. ZIEMER: Well, okay, but let me tell you --21 let me tell you, in defense of NIOSH -- and 22 again, I don't work for NIOSH -- that they have 23 in process something like 20,000 individual 24 claims. They have -- they have done dose 25 reconstructions on a little more than half of

1	those already, and obviously not everybody is
2	successful in their claim, as you might expect.
3	Some are turned down and some are not. But in
4	any event, they do
5	MS. HOLT: (Off microphone) Why don't they say
6	(unintelligible)
7	DR. ZIEMER: they do have
8	MS. HOLT: (unintelligible), why don't they
9	say (on microphone) we do not know whether this
10	man was exposed? We know that this man was
11	exposed. We know that this man died at the age
12	of 24, after he was exposed. What else do you
13	need?
14	DR. ZIEMER: Our Congressmen have put in place
15	a law which mandates certain steps that we must
16	follow legally. We cannot simply say someone
17	worked at Oak Ridge and therefore they are
18	entitled to this. That's not the way
19	MS. HOLT: I thought the President
20	DR. ZIEMER: the law is written.
21	MS. HOLT: I thought President Reagan put that
22	into effect.
23	DR. ZIEMER: Well
24	MS. HOLT: (Off microphone) We don't need to be
25	here (unintelligible) Washington. Right?

1 DR. ZIEMER: Sure. Yeah, understood. We --2 MS. HOLT: (Off microphone) (Unintelligible) be 3 there when the (unintelligible) --4 DR. ZIEMER: We -- sure. We -- this group is 5 doing the best it can to carry out what we are 6 legally required to do, as is NIOSH. 7 MS. HOLT: Okay, then why are --8 DR. ZIEMER: We understand --9 MS. HOLT: Why are you here tonight then? 10 are you here for? 11 UNIDENTIFIED: To listen. 12 UNIDENTIFIED: To listen. 13 DR. ZIEMER: We are here to listen tonight, 14 insofar as we are -- our responsibility -- you 15 perhaps weren't here when we talked earlier, 16 but the responsibility of this Board is to 17 review what NIOSH is doing in dose 18 reconstruction --19 MS. HOLT: We all know what they're doing. 20 DR. ZIEMER: -- and inso--21 MS. HOLT: Why send you all here to tell us? 22 DR. ZIEMER: It -- we're here to advise the 23 Secretary of Health and Human Services, but in 24 the process, we do -- we do like to get 25 information from people such as yourself, which

1 -- which points out -- and all of this -- all 2 of this information is -- goes into the public 3 record that points out the frustrations that 4 many of you feel. That's an important 5 component. We need to -- we need to make that 6 information known, in some cases to Congressmen 'cause they are listening, too, and they --7 8 they know these frustrations. And if the law -9 10 MS. HOLT: Evidently no one --11 DR. ZIEMER: -- if the laws need to be changed 12 13 MS. HOLT: -- no one is listening. DR. ZIEMER: Well, we're -- we're hopeful that 14 15 they -- that they will. And if the laws need 16 to be changed, that process, you know, can go 17 forward. But it -- it obviously is a 18 frustrating one. You know, I -- I can tell you 19 that even the Board -- we -- we share some of 20 those frustrations, trying to do what we're 21 legally required to do. But I recog-- I 22 recognize what you're saying and, you know, we 23 really will -- are trying to do what we can to 24 -- to address those issues. They are very, 25 very difficult -- very difficult, and -- and

1 we're not saying that it's easy, particularly 2 in these cases where we really don't know. 3 we don't know and can't find coworkers or 4 someone or groups that represent that person, 5 and when they do they make what are called claimant-favorable -- you may not --6 7 MS. HOLT: All of these people fell through the 8 cracks. There's nothing left. 9 DR. ZIEMER: Well, we -- yeah, we hope that 10 doesn't occur. We're trying to prevent that, 11 really, yeah, but -- thank you. 12 MS. HOLT: (Off microphone) All of you need to 13 this book I've got back here with all 14 (unintelligible) and all of us get together and 15 go to whoever and if we have a thousand or a 16 hundred signatures, we can get a lot done. And 17 (unintelligible) back here (unintelligible) 18 name and phone number down, we will all get 19 together. 20 DR. ZIEMER: Yeah. Well, you're guite right, 21 and it doesn't -- it never hurts to organize. 22 Are there others who wish to address -- yes, 23 please. 24 UNIDENTIFIED: I just want to make one 25 statement.

1 DR. ZIEMER: Yeah. 2 UNIDENTIFIED: I know a lady in this town 3 that's already been -- got her settlement. She 4 got it within three months, and this lady over 5 here said that from -- it took from 2001 to 6 2006 to get rejected. That doesn't make sense. 7 DR. ZIEMER: No, it doesn't. All right. 8 lady here, and then the gentleman. 9 MS. LONG: My name is Lindsay Long. I've got -10 11 DR. ZIEMER: I'm sorry, could you give it 12 again? MS. LONG: Lindsay Long. 13 14 DR. ZIEMER: Okay. 15 MS. LONG: And I've got a couple of questions 16 for NIOSH. Is -- when the NIOSH interviewers 17 call us and they do our interview, they want to 18 hurry you up because you're taking too long to 19 explain things to them, or you're asking them 20 too many questions. They say we've got another 21 interview here in another hour. I don't 22 appreciate being hurried on and passed on to 23 the next person. 24 When you're tak-- having your appeals hearing, 25 they're trying to push you on and hurry you up

1 because they've got somebody else in an hour. 2 I don't appreciate that, either. 3 I'd like to know why there's no written record 4 of the appeals hearing. The stenographer does 5 her work, she doesn't type anything down, she puts it on a cassette and then she hands it to 6 7 the hearing officer. She doesn't even get to 8 see what she's recorded. It goes somewhere 9 else. Why there's no official record, like 10 there would be in most courts, if this is an 11 official hearing? 12 DR. ZIEMER: I don't know the answer to that. 13 I think that you may be talking about a 14 Department of Labor hearing. Is that correct, 15 not a NIOSH -- yeah -- yes. 16 MS. LONG: I'd also like to know why, when 17 we're -- it may not be your issue, but that we 18 -- under the Freedom of Information Act, we 19 request log books and we can't get copies of 20 these records. We keep hitting brick walls, 21 even though we know where they exist, we know 22 where they are when we request them and we can 23 say where they are, but we still can't get 24 them. Why are we still hitting brick walls 25 from the Department of Energy?

1 DR. ZIEMER: I don't know the answer to that, 2 either. I do know that this Board has had 3 difficulty getting things from the Department 4 of Energy, also. 5 MS. LONG: Thank you. 6 DR. ZIEMER: Yes, sir -- come -- come to the 7 mike, please, so we can record. 8 UNIDENTIFIED: I just found out about this 9 meeting a little while ago so I'm really not 10 prepared. 11 I worked at Y-12 --12 DR. ZIEMER: What's your name, sir? 13 MR. O'NEAL: Earl O'Neal. 14 DR. ZIEMER: O'Neal? 15 MR. O'NEAL: O'Neal, O-n-e-a-l. I left in '86 16 and I went to work for nuclear power plant, so 17 my first job was down at Barrett Power Plant in 18 Georgia. And they asked me if I ever worked in 19 nuclear and I told them yes, I worked at Y-12. 20 And they said what's your dose rate? I said I 21 have no idea, they never told me. And so he said -- this is the NRC guy -- now he says I'll 22 23 find out. I said good -- good luck. I said 24 nobody else can find nothing out up there. So

I seen him a couple weeks later -- I already

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went to work and everything -- and I asked him, I said now, what -- what's my dose rate? He said I don't know. I said why not? He said they won't give it to us. And I said well, I thought you said you was NRC, that you could get anything. And he said well, we thought we could, but he said we -- we can't and he said we -- we're going to start you with a zero dose rate. And so I worked 25 nuclear power plants so I still -- I got dose now from that, plus what I got over there that they wouldn't even tell me about. But that's who you're working against.

DR. ZIEMER: Yeah.

MR. O'NEAL: Is people that knows how to shuffle things and hide things. So I just wanted to get in my piece -- and I been fighting them since '87.

DR. ZIEMER: Okay, thank you. Yes, ma'am.

MS. BOINET: My name's Diane Boinet. My
brother, Maurice Anthony Fitzpatrick, died
January the 25th, 2001. He worked at Y-12
plant for 27 years. I think the last time -last job he had out there he was a expediter
and he died with cancer. Okay. He had never

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been married before, no kids or anything. mother would have benefit from the program, but she died December the 18th, 2004, and I got a letter from NIOSH in January the 25th, 2005 that he had a 50 percent greater that he had received his cancer from the plant, and that's what I'm saying. It took so long, my mother was eligible for it but she died, but the program been such a hold-up and, you know, people are dying off and my brother died a awful death but no one will receive any money because he had -- he's never been married and no kids at all. And you know, I just -- it's just sad that he had to die this way and nobody would be able to get the money. But his -- his cancer -- they did say it was 50 percent greater that he received his -- his cancer from working at Y-12 plant. I'm just saying, you know, and I know how some other people feel, too, about the delay and stuff, people dying off.

DR. ZIEMER: Right.

MS. BOINET: Okay, thank you.

DR. ZIEMER: Thank you. Yes, ma'am.

MS. MOODY: I'm Shirley Moody, and I'm here for

1 my husband, Earl Moody, who passed away April 2 the 29th, 2001. He worked at Y-12 almost 20 3 years. He died of colon cancer. I have been 4 denied. He was in maintenance, which he worked 5 in nearly every plant, every building at Y-12 -6 - around the ponds repairing fences, steam 7 plants, rad houses. And I got a letter last 8 week, they've reopened my claim, but they want 9 me to have medical records from his doctors 10 saying that his cancer was caused by radiation, 11 which no doctor -- they have told me that they 12 cannot prove that. So what do I do after the -13 - now? 14 DR. ZIEMER: Okay. Again, you probably need to 15 get Ms. Moody with one of our claim people. 16 think -- are they in the corridor still? 17 get some -- we'll get some information for you 18 on next steps for you. 19 MS. MOODY: Okay, thank you. 20 DR. ZIEMER: Yeah, thank you. Who's back 21 there? 22 (Pause) Stu Hinnefeld, can you direct Ms. Moody here? 23 24 Ms. Moody, look behind you there. See Mr. 25 Hinnefeld there? Get with him and get -- she

needs some information.

MR. HINNEFELD: Okay.

DR. ZIEMER: Yeah. Okay, sir?

MR. BROWN: I'm Dennis Brown. I'm representing -- for my mother on behalf of my father. died back in 1980, worked at K-25 30-some-odd years. I was there when my dad took his last breath in 1980, you know. He was -- many months at Four Centers* Hospital. Me and my brother in college, two younger siblings at home, spent a lot of days and nights at the hospital. We'd rotate nights. My mother would stay during the day while we'd go work out for college, but like a lot of these people -folks here today, percentage shouldn't matter. One percent is too much, you know. These folks that have 45 percent, 40 percent, whatever, this -- this shouldn't be a case scenario whatsoever. I know you have guidelines and everything to go by, but again, he died of liver cancer -- covered with liver cancer. Thirty-some-odd years, you're talking about -and my dad wasn't nobody big. He was a bluecollar worker, but he loved what he did. A lot of people loved him, you know. He was there

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1 every day, very punctual, very reliable, and he 2 was -- he was -- like I said, he was a janitor 3 supervisor, you know, with five kids to raise. 4 My point to you guys is, or my question is, I'm 5 not too happy with your interactions -- you know, this is what I get from y'all for the 6 7 last three years. I could probably cover Oak 8 Ridge with the paperwork that I have. It seems 9 like every -- every paper, they come up with a 10 different scenario, something different every 11 time. Like the lady said a while ago, the 12 thing keeps dragging on and on and on. 13 My mother -- she wasn't concerned with this. don't know what -- what year did this start, 14 15 the re-- reconstruction dose, what year was 16 that you guys --17 DR. ZIEMER: It really started in 2001 is when 18 NIOSH got underway. I think the law -- the law 19 went into place earlier, but it wasn't --20 MR. BROWN: And her thoughts to us --21 DR. ZIEMER: Yeah. 22 MR. BROWN: -- we tried to get her to -- you 23 know, to go ahead and buy into this thing. 24 It's not going to bring him back, that's what 25 she told us.

DR. ZIEMER: Yeah.

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MR. BROWN: You know, it's not going to bring him -- so she wasn't too interested in -- into dollars or whatever. It wasn't till -- we was a late bloomer into this thing. It's been two years ongoing, two or three years ongoing. was then a pastor talked her into coming up here to try to do something about it. But what I'm saying is, guys, it's a small price to pay. My dad was 51 years old when he died, 30-something years at the plant -- 30some-odd years plus. I remember when I was a freshman in college, I got his watch for 30 years of business, you know, at the plant and everything. But 15 years of his life, the rest of it was spent in Oak Ridge. You know, 51 years old is how old he was when he died in So I think y'all need to be -- instead of the paperwork, I think you need more interaction, a lot of one-on-one with people that does have these claims. And you know, let them know that you care. This paperwork don't mean nothing to me. And I think I'm speaking

DR. ZIEMER: Sure, understood. Uh-huh.

for all these folks, you know.

1 MR. BROWN: If you try to interact with us, 2 we'll -- everything'll be a lot better. It's a 3 small price to pay for 30-something years of 4 service. 5 DR. ZIEMER: Yes, sir. Thanks. Yes, ma'am, 6 sure. 7 MS. MCKEETHAN: Hi, I'm Diane McKeethan*. I'm 8 pretty sick, and I was pretty sick about 12 9 years ago when I worked. And before the Act 10 was ever signed I'm pretty sure I was -- I had 11 a dose, okay. I don't know what it is, but I 12 did inventory, so I know I -- I was hot. 13 Anyway, I want to know what we, as a group, can 14 do to help you guys, 'cause you're people, too. 15 And I know you see the human suffering. 16 DR. ZIEMER: Yes. 17 And I just wonder what we can MS. MCKEETHAN: do, as a group, because sometimes it takes the 18 19 power of the people to get behind you. 20 DR. ZIEMER: Right. 21 MS. MCKEETHAN: So please tell us --22 DR. ZIEMER: Yes. 23 MS. MCKEETHAN: -- please. 24 DR. ZIEMER: One -- one starting point of 25 course is sharing what -- as you have tonight.

1 That is in fact helpful for the process. 2 may seem frustrating, but it does have an 3 impact. It has an impact on the actions of 4 this Board. It has an impact on the federal 5 staff people who operate the programs. They're 6 not all cold-hearted, really. And it does have 7 an impact I think on your legislators, who 8 really determined the ground rules on which we 9 operate. So that's a starting point. 10 Some groups do find it helpful to organize in 11 certain ways, particularly if there's some 12 political aspects that you need to -- to 13 address. But I -- you know, I can't -- I don't 14 want to get into that aspect myself, but you 15 know, what you're doing already is helpful to 16 us and we appreciate it. 17 UNIDENTIFIED: (Off microphone) 18 (Unintelligible) like a petition 19 (unintelligible) petition (unintelligible) or 20 (unintelligible) organization? 21 DR. ZIEMER: Well, of course there already is a Y-12 petition that's under review, so that --22 23 that is in process right now. What you're --24 what you're doing here -- this information gets 25 shared and actually gets tracked to see -- for

example, issues that are being raised about paperwork and so on, the -- the agency and its contractor actually track this information and try to determine how to address it, so that's helpful, as well. And sir, I think you indicated you wished to address the group. MR. SCOTT: My name's Frank Scott. I'm the

mr. scott: My name's Frank Scott. I'm the president of Local 900 in Oak Ridge. I'm working on my 30th year at the Y-12 plant. I was a chemical operator for 16 years in the 9-212 area. I'm happy to report, as far as I know, I'm in great health right now, so I'm not here to talk about my health.

But the dose reconstruction -- I have been involved in very -- probably more accidents in the Y-12 plant than most people have when I was in chemical operations, and some pretty serious things went on. And I can remember a fire that I was involved in where myself and some coworkers were involved in a chip fire where the whole room was full of contaminated smoke, and one of those coworkers today I understand has some very serious health problems. So you know, I expect somewhere down the road I may

have some, too.

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But this dose reconstruction, to think that you guys can figure out how much I've had since I've been out there, no way. No way. Do I have confidence that the folks out there have -- have gave you accurate records, even the ones that you do get? No way. No way. I have what is considered now probably as a -as -- if -- the people out there, when you get your dose record every year, most of them is going to read zero. Mine don't read zero. Mine reads around 300 or so, which -- which even to you all, that ain't a big number. But I also was involved in a program back in the mid-'80s -- I was a volunteer for RadCon, believe it or not, to -- their -- their effort was to prove that there was no insoluble uranium in the Y-12 plant so we could do less testing on people for contamination purposes -urinalysis, as a matter of fact. And I regret that I was a part of that study because what it resulted in is -- is they went from a monthly urinalysis to -- to doing a urinalysis whenever you went to a rad area. And shortly after that, I was a representative of the chemical

1 operators at that time, and shortly after that 2 I had a man come up with unexplained Y material 3 in him, which is what we just proved we didn't 4 have any in Y-12. 5 So do I have any faith in what's going on out there? No. 6 No. 7 Do I feel like that we've had skewed records 8 because of people trying to make sure that --9 that the folks look good that's in charge of 10 the plant? Yes. 11 If I'm a contractor and I'm out there and I get 12 numbers coming at -- going at DOE saying we've 13 got this person radiated this much, this much, 14 this much, and then I've got to turn around and 15 the folks that work -- that work for me and 16 their pay raises -- you know, I'm -- I'm 17 responsible for pay raises, they're going to --18 they're going to skew numbers, and they do, and 19 they have, and they will. 20 My -- my saying is you will -- you will not get 21 an accurate reading. I also say if anybody's 22 really interested in what's going on with our 23 health out there, that we'll separate that and 24 put that under the Department of Health, get it 25 out from under the money that -- that's -- that

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it -- then it probably will help that issue. Thank you very much.

DR. ZIEMER: Okay. Thank you. Yes, ma'am, uh-huh.

(Unintelligible) microphones, but I'm Valerie Maner and this is my father, Ralph Delozier. I am a medical technologist and I specialized in nuclear medicine. I deal with radiation badges every single month. I -- we haven't dealt with this reconstruction thing because we're not there yet. There's no way you can take -- like he was an engineer. He would go out in the plant. You can't take another engineer and make it the same. They don't do the same things. My heart goes out to these people 'cause you cannot reconstruct an individual's radiation. You can't do it. You cannot do that, and you know that. You're a physicist. Right? You know you can't do that. I know you can't do that. It's not fair.

DR. ZIEMER: Thank you. Ma'am, did you -UNIDENTIFIED: I just want to ask a question.

DR. ZIEMER: Would you approach the mike again? Sorry to make you squeeze out of there, but we do need to get your name and be able to hear.

1 MS. FOSTER: I'm Ellen Foster. I'm here on 2 behalf of my father. He was turned down after 3 fighting for three years, and I got a letter in 4 December and it said I may have -- I may 5 request a reconsideration. It says such a 6 request must be -- must be in writing and must 7 be made within 30 days of issuance of this 8 decision. It was clearly -- it must clearly 9 state the grounds upon which reconsideration is 10 being requested. The request for re-- for 11 reconsideration should be sent to the 12 Department of Labor. 13 If there's anyone here tonight from the 14 Department of Labor, I would like to talk to 15 them. 16 DR. ZIEMER: Sure, there are Department of 17 Labor folks here. They may be in the corridor. 18 DR. NETON: Out in the hall. 19 DR. ZIEMER: In the hallway? Jim, can you 20 direct Ms. -- is it Foster -- to someone from 21 Labor --22 MS. FOSTER: Also, and employee of NIOSH told 23 me the last time I talked to her that from 1959 24 until 1961 they did not wear badges at Y-12, 25 and my father was there. So he died of cancer,

but I've been denied after three years. I worked there in the guard department. I was injured and I -- I got hurt and they -- and they say, you know, I had a body count. I never had a body count and I was all over the plant. I crawled around the attics, I crawled around everywhere, in -- in the buildings that they said was really hot, but I never had a body count. I never had any kind of -- of count. When I left there I didn't even have a physical, so --

DR. ZIEMER: What years were you there?

MS. FOSTER: I was there from 1979 to 1990.

DR. ZIEMER: From 1979.

MS. FOSTER: And I didn't have a physical when I left there so they don't know what I left out of there with. So -- but I ha-- I do have thyroid problems. I'm now a diabetic, and my husband was there during the -- during the war. He -- he was a -- he was a chemist. He was supervisor out there, and now he's got Alheimer's (sic) and he's also lost his hearing. They say they don't pay for the hearing loss, but we spent four hours in Knoxville when we got this letter saying we

should have the examination. Well, they said I didn't have anything, but I do have diabetes and I have -- I have a thyroid problem, but I didn't have it until I left there. But what is going to become of this -- of this thing -- this examination where they took it over in Knoxville from ATLC? Can anyone give me an answer?

DR. ZIEMER: I don't know, but if you -- if you raise that issue with -- Mr. (sic) Neton can perhaps direct you to where to find the answer -- yes. We'll -- we'll try to help find what you need.

MS. FOSTER: Thank you.

DR. ZIEMER: Yeah. Yes, ma'am.

MS. ALLEN: I'm Janice Allen and I'm acting on behalf of my mother, Nancy Thomas. We -- my mother was diagnosed with breast cancer and she had 14 cancerous tumors in her lymph nodes. They had to be removed in 2000 and we applied in 2001 and they turned us down in 2004. Well, since then she's done -- from the breast cancer and the cancerous lymph nodes, it's done spread. She has liver cancer, she has bone cancer, it's in her brain and everything.

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Well, they turned us down and I called and asked them, you know -- you know, that's the reason why it's -- spreads, it's from the breast cancer and the lymph nodes cancer, and they said well, they doubt very seriously if I could reapply because that's a different case. So what are you supposed to do if they turn you down, they think -- I mean cancer is cancer. Once you got it and it spread throughout your body and -- but they're saying like well, you know, that's -- that didn't happen 'cause of that. And they said because of the breast cancer, because the lymph node cancer, it wasn't enough doses, so now that it's all through her body and she's on hospice and all that, they're saying like well, it's nothing -no big deal, really, is what they're saying. I don't think we know the answer DR. ZIEMER: to that here at the Board table, but on specific cases like that, again, you need to get with one of the caseworkers and have them follow up to see if there is an opportunity for that to be considered.

MS. ALLEN: Thank you.

DR. ZIEMER: Yes. Sir, another comment?

1 MR. DISHMAN: Could I --2 DR. ZIEMER: Yes. 3 MR. DISHMAN: Could I ask the question of who -4 - is it Energy or Labor that is abusing these 5 people on this reconstruction? Is it Energy 6 Department or is it the Labor Department? 7 know --8 DR. ZIEMER: You'll put me on the spot. 9 MR. DISHMAN: -- you see, that's the problem. 10 DR. ZIEMER: I don't -- I don't know the --11 MR. DISHMAN: It's to --12 DR. ZIEMER: -- answer to that, I --13 MR. DISHMAN: -- keep the hat away from these 14 people --15 DR. ZIEMER: No, no. 16 MR. DISHMAN: -- you know. 17 DR. ZIEMER: We're -- we're trying --18 MR. DISHMAN: It's hide the hat. 19 DR. ZIEMER: We are trying to find the 20 information that can be used for this --21 MR. DISHMAN: But surely you know if we're --22 when they protest, and they protest too little, 23 should they protest to the Labor Department or 24 the Energy Department? We hope the Labor 25 Department cares more about them than the

1	Energy Department because they've been down
2	that road and there's big ol' ruts running in
3	that road.
4	DR. ZIEMER: Yeah, yeah, I understand. I don't
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6	MR. DISHMAN: But where does the buck stop?
7	DR. ZIEMER: Yeah, I don't
8	MR. DISHMAN: The buck's got to stop somewhere.
9	DR. ZIEMER: Yeah, I don't know the answer to
10	that. We're we're trying to address what we
11	can. We we've heard many concerns here
12	tonight. We will you know, we're trying to
13	address those. I don't know how effective we
14	will be, but we will try. That's all I can do
15	tell you tonight, you know.
16	MR. DISHMAN: But we but we don't know what
17	Department we're
18	DR. ZIEMER: And I'm
19	MR. DISHMAN: having problems with.
20	DR. ZIEMER: I'm not sure. You know, if I
21	said one or the other
22	MR. DISHMAN: Does some of our Board members
23	know?
24	DR. ZIEMER: No. Well, you know, it's like
25	who's to blame for I mean

1 MR. DISHMAN: If you can't figure out who to 2 blame --3 DR. ZIEMER: Well, you know, the --4 MR. DISHMAN: -- you can't get results. 5 DR. ZIEMER: The problem is a complex problem 6 that has grown up over the years. I mean --7 MR. DISHMAN: Well, we agree. 8 DR. ZIEMER: -- all of you folks -- all of you 9 folks -- and us, most of us have had nuclear 10 experience -- we -- we entered -- we entered 11 these fields really, in a sense, on behalf of 12 our country. All of you were, in a sense, 13 volunteering, in many cases. You now see that 14 you didn't know fully perhaps what the risks 15 were that you were facing, and -- and I'm not 16 even sure the agencies at that time knew those 17 themselves, and that's probably part of the problem. They didn't monitor appropriately, by 18 19 today's standards. And we're going back and 20 trying to correct mistakes of the past, and 21 it's very difficult to do, very difficult to --22 MR. DISHMAN: Please let the record show that 23 no one knew what Department this falls under. 24 DR. ZIEMER: Well, we know who has certain 25 responsibilities. You asked who's to blame for

1 the problems --2 MR. DISHMAN: Well, who has responsibility for 3 the dose reconstruction? 4 DR. ZIEMER: NIOSH is responsible for 5 conducting dose reconstructions. MR. DISHMAN: Is that under Labor? 6 7 DR. ZIEMER: No, that's under Health and Human 8 Services. 9 MR. DISHMAN: Health and Human Services. 10 DR. ZIEMER: Right. 11 MR. DISHMAN: It's not under --12 DR. ZIEMER: Labor -- Labor's responsible for verifying certain things -- the medical 13 14 records, the employment records and doing the 15 determination of probability of causation. 16 those responsibilities are split. Department 17 of Energy has the responsibility for providing 18 records, and we're aware of your concern about 19 the records. So it's split up. 20 MR. DISHMAN: And that's why we can't ever --21 DR. ZIEMER: We understand. 22 MR. DISHMAN: Thank you. 23 DR. ZIEMER: A gentleman over here on my left. 24 MR. RUSSELL: My name is M. L. Russell. My 25 previous badge number's 29562. I'm a sick

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worker. I was one of the original ten that signed up on this program. That was back in 2000, signed up again in 2001, still haven't gotten any results yet. That's been five years. I had Drs. Lockey and Byrd sign that my exposures came from work. It was supposed to have put me on through. Still haven't heard anything from it.

Everything that we've given -- I've given repeat documents time and time again, and a lot of people wonder how they've been exposed to things. That right there is DOE's own documents. That's Rifle Range UF-6 explosion test, which people can see just the things that went on in your back yard that you didn't even know about, just showing that there was a little test that went on that people didn't know. And such things as the cooling towers that people drove through in the mornings that the fog was so thick that they couldn't see where they was going. If you go back and you check your calculations on what they checked just for the chromium on the towers, you'll find that they calculated that incorrectly. They calculated it for one tower, not out of

1 all of them. 2 It's your own DOE documents that everybody has 3 that they know it's there, but yet when people 4 comes to them with them that I've presented it 5 time and time again -- cross-water connections, I was on a committee for it. All of a sudden, 6 7 the funding for it ended. It never got 8 addressed. It was supposed to been handled by 9 an independent agent. It got shut down. 10 Time and time again we've showed that the 11 problem exists, but it's another panel, a 12 different panel, different people, more sick 13 workers. Information just gets recorded and 14 don't get handed down. I've had several 15 surgeries. I hope I make it past all of them, 16 but these people are just getting the runaround 17 like everybody has. Look at the documentation 18 that people's using you. Let's see some 19 results. 20 DR. ZIEMER: Thank you. Yes, ma'am. 21 MS. THOMPSON: I'm sorry, I just have two more 22 comments. Number one --23 DR. ZIEMER: Give your name again --24 MS. THOMPSON: Oh, Dorothy Thompson. I'm told

that you cannot question NIOSH's dose

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1 reconstruction. Is that correct? If you go to 2 an appeal --3 DR. ZIEMER: I think you can always question 4 (unintelligible) --MS. THOMPSON: No, if you go to an appeal, you 5 6 cannot question NIOSH's dose reconstruction. 7 DR. ZIEMER: Oh, the -- yeah, the appeal -- the 8 appeals are through the Labor Department, so 9 Labor --10 MS. THOMPSON: Is that right --11 DR. ZIEMER: Yes. 12 MS. THOMPSON: -- Dr. Wade? Even though we 13 know that their estimates -- that they're 14 invalid, that they're inaccurate and that 15 they're only estimates, we can't question them. 16 That's pretty presumptuous. 17 Then secondly, it's my understanding, Mr. -- is 18 it Dishman? -- that we are to blame our own 19 state senators for not joining the original bill -- and correct me if I'm wrong -- that put 20 21 Y-12 into the group that could claim these 22 things without dose reconstruction. Is that 23 right? 24 DR. ZIEMER: I -- I'm not --25 MS. THOMPSON: The Kentucky senators, the --

1	DR. ZIEMER: I'm not suggesting
2	MS. THOMPSON: Ohio senators
3	DR. ZIEMER: you blame your state senators
4	for what they may or may not have done
5	MS. THOMPSON: Well, it may have been an
6	oversight at the time, but at any rate, Y-12
7	was not put into the original document tell
8	me what the that's true, that it was not put
9	in as a special group like Paducah and
10	Portsmouth.
11	DR. ZIEMER: Right.
12	MS. THOMPSON: Okay. So that's who we really
13	should
14	DR. ZIEMER: And I I don't
15	MS. THOMPSON: be after.
16	DR. ZIEMER: think this Board really knows
17	the workings of Congress in that case, what
18	what factors went into the original
19	determination of who was or wasn't in the
20	MS. THOMPSON: It's my understanding
21	DR. ZIEMER: cohort.
22	MS. THOMPSON: that the Tennessee senators
23	were not in there fighting for Y-12.
24	DR. ZIEMER: I don't know the answer to that.
25	MS. THOMPSON: Are any of the representatives

1 here tonight? 2 UNIDENTIFIED: Of course not. 3 DR. ZIEMER: Yes, sir. 4 UNIDENTIFIED: (Off microphone) 5 (Unintelligible) 6 DR. ZIEMER: Mr. Elliott from NIOSH --7 MR. ELLIOTT: Let me answer your earlier 8 question, though. Once you receive a dose 9 reconstruction from NIOSH, you are asked 10 whether or not you have any additional 11 information or not to provide. You're asked to 12 fill out -- sign the OCAS-1 form, and then it goes over to Department of Labor. You can 13 14 appeal on the dose reconstruction as to whether 15 or not we applied the methods appropriately. 16 You can question the application of 17 methodology. You can't question the methods 18 themselves. 19 The Board is charged with evaluating our 20 methodology and working on that, so just -- I 21 wanted to answer your question about what you 22 can appeal on. Okay? 23 UNIDENTIFIED: (Off microphone) 24 (Unintelligible) 25 MR. ELLIOTT: You can appeal on whether our

1 methods were applied appropriately. You can't 2 question methodologies. 3 UNIDENTIFIED: Why? 4 MR. ELLIOTT: I'm sorry, that's just the way 5 the law is written. 6 DR. ZIEMER: Okay. Sir. 7 UNIDENTIFIED: (Off microphone) 8 (Unintelligible) 9 DR. ZIEMER: There's a law that specifies how 10 they are to do dose reconstructions, basically, 11 is what is being said. And -- and if they 12 don't -- if they don't apply that law 13 correctly, that can be --MR. ELLIOTT: We were required by law to 14 15 provide recommendations in regulations on how 16 we go about doing dose reconstructions. 17 regulations were reviewed. You can comment on 18 the regulations. But once you have a dose 19 reconstruction, the Department of Labor 20 regulations and our regulations in NIOSH only 21 allow appeal on whether or not the dose 22 reconstruction methodologies were applied 23 appropriately, not on the methodologies 24 themselves. I know it's confusing, but I

wanted to answer your question 'cause I felt

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you didn't get an answer to it earlier.

unidentified: (Off microphone)

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(Unintelligible) understand what you're talking about here.

DR. ZIEMER: Okay. Sir, go ahead.

MR. LEE: My name's Otis Lee. I passed on the first go-around. I may or may not have a dog in the fight here tonight. I've noticed some of the crowd has left, but I'd like to comment just a little bit. I retired from DOE with about 23 years. I was a courier, nuclear transport special agent, and I would like to say that this -- it just -- from listening to these horror stories, it appears that the NIOSH reconstruction is some sort of a dog going down a rabbit trail looking for a scapegoat and I don't quite see -- you cannot dispute scientific methodology and things of that. mean what's -- two and two is four. you have -- you're trying to use a reconstruction as a litmus for folks who -- a litmus test for folks who haven't even worn dosimer (sic) badges, so it's -- it's an invalid method of trying to say you -- you do or do not qualify as to -- for this 50 percent

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situation because people weren't there. So not meaning to make waves, but many folks are familiar with the Love Canal/Erin Brockovich situation, and I don't know if a class action lawsuit would be more appropriate to get power to the people or not, but I'm just saying that perhaps that may be some help. I know that situations I was involved in as a transportation security specialist, there were many times health physics folks would come out, they would find contamination and tell us all to leave, go up here, and then next thing you know, it's all clear, and we'd be setting off alarms and things of that nature. We had a staging area in the middle of a contaminated area that we would wait for shipments and things of that nature. So a lot of the -- the situations we were in were very -- just iffy, we just did not know what we were getting exposed to. So -- but I would encourage those folks to sign that little ledger that the lady has sent around and perhaps we'll all get in touch and maybe we can -- maybe we may be able to have more muscle than what you folks have. So thank you.

1	DR. ZIEMER: Okay, thank you. Yes, ma'am.
2	UNIDENTIFIED: (Off microphone)
3	(Unintelligible) I didn't get your names and
4	phone numbers. My name is Faye Holt. My phone
5	number is 865-882-5618.
6	DR. ZIEMER: Okay. Thank you. Go ahead,
7	ma'am.
8	UNIDENTIFIED: (Off microphone)
9	(Unintelligible) leave your name and number
10	(unintelligible).
11	MS. KILEY: I'll make this very brief. I'm
12	representing my father's case. His file number
13	is 420055522. His name is Clifton O'Neal.
14	DR. ZIEMER: And your name is
15	MS. KILEY: Is Debra.
16	DR. ZIEMER: Debra O'Neal?
17	MS. KILEY: Debra Kiley.
18	DR. ZIEMER: Debra Kiley.
19	MS. KILEY: Yes.
20	UNIDENTIFIED: (Off microphone) Ma'am, can you
21	stand (unintelligible) to the microphone
22	MS. KILEY: Yes, I will.
23	UNIDENTIFIED: (unintelligible) hear you?
24	MS. KILEY: Yes. Back to our dose
25	reconstruction, as usual. You know, I did some

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research on this, as you can see. I work in the medical field, and my father had metastatic adenocarcinoma of the rectum. He didn't have just a few sites, but he had five primary sites of cancer. And you know, based on data from the American Cancer Society, cancer facts and figures 2005, the probability that cancer will result from radiation exposure increases as the dose increases. And NIOSH dosimetry calculations for my father, based on their findings, the District Office calculated the probability of causation for rectal cancer and determined that the probability that the cancer was caused by exposure to radiation during employment is 10.03 percent. And he worked out there nearly 30 years as a machinist. He began employment in '54 at age 35 and in 81 he was age 63 and ten months later he was diagnosed with final stage cancer. And ten percent, hmm. When I looked back at all the information that I was given, the determination is made based upon guidelines also developed by NIOSH, and incorporated into an interactive computer program that OWCP uses to calculate the probability that a claimant's cancer was caused

by exposure to radiation. Whatever your methodology is for that, you know, we need it in laymen's terms and I agree with -- there was a young man who spoke earlier -- it's not a percentage. Any -- any amount of exposure is over-exposure, and it is not -- it's just not even feasible that this is going on. We know of a secretary at K-25 who got compensated, and my father was a machinist who -- he ground uranium, and please explain this to me and take this back to whomever it is necessary. Thank you.

DR. ZIEMER: Thank you. Yes, sir.

MR. DELOZIER: I'm Martin Delozier. This is my father over here. I didn't come prepared to talk tonight, but he worked at the plants, I also worked at the plants for ten years. Just to give you a quick history of what I did, I was -- when I first started at the plants I worked at the -- down at K-25. My job was -- when I first got there was take the Geiger counters -- they brought equipment out of the plant, to test what the radiation level was, whether it went to this yard or this yard.

But all that aside, what I'm -- want I want to say here and try to get an understanding of what we're supposed to do. We're looking for information to turn in to you guys to request or whatever, and ask for medical records which these people cannot get from the plants, very, very difficult. You've admitted yourself that you have trouble getting them. Just looking for some guidance what to do.

Also, some of the other things that you're looking for, by everybody's admittance, you just have trouble getting this information. So to help these people find out what we need to do next, we're looking for some answers from you guys. And if you don't have the answers, these people are lost. And we're looking for answers from you all. That's what these meetings are for, answers from you guys to tell us what to do, what papers to fill out, how to get the papers if we can't get them and you can't get them. And to reconstruct the dose things, as everybody knows, is impossible. I mean I worked out there and never had a dose meter.

DR. ZIEMER: Right. NIOSH has people on deck

1 here to help with individual cases such as 2 yours, so you need to --3 MR. DELOZIER: Well, it's not indi-- I'm just 4 looking for information for the whole people. 5 Tell us where to go next --DR. ZIEMER: Oh --6 7 MR. DELOZIER: -- 'cause you're requiring 8 information that we cannot get. You're 9 requiring information that can't be done. 10 DR. ZIEMER: Right. 11 MR. DELOZIER: I mean like my father had cancer 12 and the records were shredded seven years ago, 13 just like other people here. I mean we've got 14 a new instance of it now and we do now have current records. He's -- 7th of this next 15 16 month he's going for surgery. So that'll help 17 a little bit there, but we're looking for 18 general information for everybody as what steps 19 do people need to do since they're up against a 20 brick wall. 21 DR. ZIEMER: Right. 22 MR. DELOZIER: And they cannot get anything --23 cannot do anything. We're spinning our wheels 24 and going nowhere but backwards. And that's 25 what the answers we're looking for. Thank you.

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That's good.

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DR. ZIEMER: Thank you. Yes, right here, sir.

MR. HACKWORTH: My name is James Hackworth.

I'd like to commend the panel in regard to the composition of the panel. I think it's admirable that you have people from the hourly workers as well as professors and a -- and a variety of people. That's -- that's required.

But I would like to say this in -- in regard to being able to reconstruct a -- a dosage over a period of years, whether it be at K-25, Y-12, X-10, Fernald, wherever it may be. Here is some of the problems that there's been a lot of encountering -- my friends, other folks, many of them are dead, but here -- here is -- almost each and every one encounters. If you go back to 1943 when it first began, I had a brother that worked at Y-12 initially and in regard to the -- oh, golly, I can't think of the name -the initial production in Y-12. But here's what a lot of folks are encounting (sic). There was no records kept prior to 1950 at Y-12. I think a lot of these folks know that. So here's -- here is a huge question question that I've got. How can you possibly go back

and reconstruct something unless you talk to a lot of these people that have been there? Many of these people are dead. You cannot go back and give those people a name to come back and talk with. Okay? So therefore, I would -- I would like for the gentleman to explain how they reconstruct with no records.

Another point I'd like to make perhaps some of the folks are not aware of, the dose rates have changed over the years. I started working in 1951. The allowable dosage rate was much higher then than it is today. Now there's another thing that should be taken into consideration, and -- and I hope -- I hope it is. But if it hasn't been, it needs to be implemented.

So, you know, here these people say no dosimeters, no -- everyone had a badge -- film badge of some sort, but I -- I would agree there's -- there's many, many people that did not wear dosimeters that should have been.

Another problem area that we have in regard to being able to produce adequate records for their defense -- or not their defense, for their record or their loved ones' record -- is

the fact that I do not feel that the contractors got -- they should not be allowed to get away with not some involvement in this. Okay? Because they had the responsibility to see that these industrial hygienes, health physics and the other programs to protect the workers were carried out. They did an inadequate job. There is no question about it. So therefore, the burden is coming back to the individuals to -- to prove something that's -- that's impossible to prove.

Now I understand the gentleman and his calculations and methodology, the whole bit.

But there's a big element missing -- no records, 1943 to 1950. So pray tell me, how can you go back and calculate something to non-existent records? They do not know the buildings he worked in, the level of activities these individuals were working, and the dosage rate they were in. Huge problem, gentlemen.

DR. ZIEMER: Well, let me give you the quick answer. If there are no records, no monitoring records, no dosimetry records and no records of what sources were present, then dose cannot be reconstructed, and that would be the basis for

1	a Special Exposure Cohort, though.
2	MR. HACKWORTH: Yet these folks yet they
3	died of the various type cancers.
4	DR. ZIEMER: Right.
5	MR. HACKWORTH: Okay?
6	DR. ZIEMER: So there and there very well
7	may be groups, depending on the years and the
8	locations, where that is the case. And if that
9	is the case
10	MR. HACKWORTH: That is very much the case.
11	DR. ZIEMER: Yeah, right.
12	MR. HACKWORTH: Many of those folks are
13	deceased.
14	I would like to just one (unintelligible)
15	one final message. Okay?
16	DR. ZIEMER: Okay.
17	MR. HACKWORTH: Mr. (sic) Wade, I understand
18	that you you run back and reporting to Human
19	Services. Is that that's your
20	responsibility. Is that correct?
21	DR. WADE: Correct.
22	MR. HACKWORTH: Okay. You heard the you
23	heard the voices of the people tonight. You
24	heard other meetings, perhaps. It's I'm
25	going to kind of quote a little bit from

1 another person that made a statement one time. You heard the message. You heard the people 2 3 and their statements. Their statements do fit, 4 so you should go back and tell your folks to 5 submit. 6 DR. ZIEMER: Okay. Thank you. 7 MR. HACKWORTH: Thank you. 8 DR. ZIEMER: The hour's growing late and I see 9 many people are leaving. I feel like perhaps 10 we should officially come to a close -- well, 11 we have one --12 **UNIDENTIFIED:** May I say something? 13 DR. ZIEMER: You certainly may. Please come to 14 the mike. I don't want to call it off too 15 soon, but I know many folks are leaving and 16 that indicates --17 MR. BOWERS: I'm Leonard Bowers. I spent 44 18 years at Y-12. Most of these people out here 19 I've worked with. I'm now 76 years old. 20 came to Oak Ridge High School when I was 14 21 years old. This creek here -- creek behind 22 this building, I used to play in it when I was 23 a child. Then when I went to work in Y-12 in -24 - in 1950, I saw what went in that creek, and I 25 used to wade there and swim in this creek out

here. And then finally they build a settlement pond out there. But this gentleman that just spoke a few minutes ago, he was trying to think of the Tennessee Eastman Corporation. Now they're the people that had the records back then. They're out of Kingsport.

Now I'm an old-timer out here, and I still have my memory. But what these people have gone through with -- I worked in the mercury, I waded in it. I've left out there when my shoes were so hot that they would -- that they'd take them up and we'd wear little soft shoes to the change-out, and they disregarded safety altogether.

When the tiger team came in from Washington, you know, they shut the plant down out there. And one of these gentlemen came in my lab -- by the way, I made printed circuit boards and I've dealt with gold, I've dealt with platinum, you name it, I worked in Y-12 from one end to the other. I went to work there in 1950 and that's when we was taking the silver out of the tracks. I was a crane operator. I served my apprenticeship out there, and I left and I went into the military and I spent four years in the

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Air Force and came back. And while I was gone, my raises went on, my seniority went on. But we'll get back to some of the things that these people have lived with. Let me tell you, right now -- I just found out recently -- I went to the Welmouth School. I didn't come prepared to speak to night or anything. But I have just got a report back and the lady, she was out of Nashville that did these tests on me, and when she first looked at my fingers she said Mr. Bowers, she said you've got heavy metals in your body. Well, right out in the car right now I've got -- I've got the stuff out there, and the rating -- like arsenic, beryllium, mercury, sodium, potassium -- I worked with all that stuff. And I start tomorrow for tests to find out about these metals that's in my body.

But I know what these people went through. I was there. I spent six and a half years on the Brigger* reactor, until President Carter shut us down. I spent 12 and a half years in biology. Now I've been around some very smart people in this world, and I went all over that plant from building to building, and I've got

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pictures of things that I worked with. We worked with asbestos and back in the very beginning we worked with carbon tet. That's what we cleaned electrical parts with.

DR. ZIEMER: Sure.

MR. BOWERS: When the tiger team came in, do you know they shut my lab down, and the gentleman -- he left me his card; I don't know if I still have it or not at the house -- but they were giving me a hard time in that plant, and he set down and he talked to me, and I told him -- well, what they did, they condemned that lab. When you process printed circuit boards, you gold plate, and that gives off phosgene gas. The roof -- there was one man, he came in my lab and he said Leonard, he said I was up there at the bus stop going to the cafeteria and he said I looked up there on that roof, he said I could see it, there's a big hole in your roof. I said what? He said there's no exhaust. Now I won't go into details, it's late and a lot of you people have talked, but I've been from one end of that plant to the other. I've worked in every building out there. I've worked with some of the smartest

1 scientists in the world.

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And it's been the educational -- I've put my hands down in centrifuge that were so hot that -- that they would leave with it. Well, I'll give you example. Back -- let's see, this would have been 1959, was working 9201-2, foreman. I asked him for 200 amp disconnect switch. Well, he was a retired colonel, and I asked him when he came up to me and I told him what I needed. I thought he'd order me a switch. Well, he comes wagging one back and I said -- well, his last name started with a K, I'll just call him Mr. K. -- I said Mr. K., where did you get this switch? Back on the mezzanine. And I said has health physics checked it? He says are they supposed to? said yes, and so we call health physics in and a fella -- his initials were D.W., he's dead now -- and they put him on to wash that switch. Now this was a switch we was going to put on out in the hall. So what does he do? When the health physics man gets down there, he puts his Geiger counter on it and Mr. -- well, Mr. W., I'll call him -- he washed that thing for four hours, and when the health physics man came

1 back he said send it to the burial grounds. 2 And another thing I can tell you that went on 3 out there -- I'm an old-timer, and I'd like to 4 share this with you. 5 DR. ZIEMER: Okay. Remember now, folks are 6 getting tired. 7 MR. BOWERS: No, this is -- this is the last. 8 In 19-- this would have been 1955. Many of 9 y'all know Charlie Robertson, don't you, that 10 I was Charlie's apprentice. We worked died. 11 together over a year, and we went in 9201-1 and 12 we were on the second floor and here I am, I 13 just got back from the service, and the guys 14 were talking that they had connected the 15 thorium line to the drinking water fountain. 16 Yes. Now I don't know if it was actually 17 turned on or not. We just said let's get out 18 of here. 19 So I didn't come prepared to talk tonight, but 20 like I say, I go way back. And when -- I was a 21 usher at the Grove Theater back in the '40s, 22 and that's my picture in the paper y'all see 23 which says "The Atomic Bomb, the Beginning or 24 the End?" That's me when I was 16 years old, 25 and I know what has gone on. And I lose my

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trend (sic) of thought sometimes, but it's been interesting. I love living in Oak Ridge. I graduated in 1947, and I've been around quite a bit. But these people here, I don't know what -- if I have problems or not, but you name it, I know more about Oak Ridge than any of y'all in this room because when you was an electrician in Y-12 you would go from one end to the other, that plant. You never knew where you were going on overtime. And these people are hurting. I've worked with them -- well, you spoke of Jack Case a little while ago. Well, Wayne Wallace was his first -- was his first wife, and Wayne -- I was working with him, and he developed this problem down in Alpha 5 in the mercury. And I felt sorry -she did get a settlement, I believe. But like I say, I worked with many of them out there. I worked with Herman Postman, how many of y'all know Herman? Well, I started with Herman back when he first came here. He was 26 years old and he worked in 9204-3, and I went down and hooked up his vacuum pumps and I made a prediction. I said there's a young man downstairs that's going to go up in this

company, and he went to the top. He became a vice president.

But we had a lot of things we shared together. We worked on the DCX program. You name it, I worked on it. Every time they'd run out of money, I'd go somewhere else, and these people -- I worked with some of the finest people in this world. And these people right here, I've worked with them. I've changed the motors out for them, their lights and all. But I'm just proud to be an American. And what I like most about when I was out there, many of them dreaded going -- taking a lie detector test, known as the polygraph test. I thought it was a honor to go up there and take that polygraph test, that I was a red-blooded American and I was proud of this country, and that's when I went in the service and then I saw things change. And the man who was head of security out there and he was head of all the quard department, and I used to sit in his lap when I was a little boy. He lived with us at Tellico Plains, and I would -- he would tell these stories. But he was one of the finest men -well, he is dead now, but I didn't mean to talk

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1 so long, but I just came in here and just heard 2 these people, and I know how they're hurting. 3 And I appreciate any help that y'all can give 4 them. 5 DR. ZIEMER: Sounds like a good note to end our 6 evening on. Thank you very much for sharing 7 that. Thank all of you for coming tonight and 8 sharing with the Board. We appreciate it. 9 We will be meeting again all day tomorrow. You're welcome to return and learn more about 10 11 the dose reconstruction process. 12 (Whereupon, the meeting was adjourned at 9:10 13 p.m.) 14

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CERTIFICATE OF COURT REPORTER

STATE OF GEORGIA COUNTY OF FULTON

I, Steven Ray Green, Certified Merit Court Reporter, do hereby certify that I reported the above and foregoing on the day of January 25, 2006; and it is a true and accurate transcript of the testimony captioned herein.

I further certify that I am neither kin nor counsel to any of the parties herein, nor have any interest in the cause named herein.

WITNESS my hand and official seal this the $7 \, \text{th}$ day of March, 2006.

STEVEN RAY GREEN, CCR

CERTIFIED MERIT COURT REPORTER

CERTIFICATE NUMBER: A-2102