BURKHOLDERIA SPP. INFECTION CASE INVESTIGATION FORM

Instructions

Please complete as much of the form as possible. The instructions below explain each variable. If you have questions, please contact Bacterial Special Pathogens Branch at (404) 639-1711 or <u>bspb@cdc.gov</u>.

Send the completed form with all personal identifiers removed to CDC either by:

Email: bspb@cdc.gov

Fax: (404) 929-1590

DCIPHER: contact <u>bspb@cdc.gov</u> for more information

Reporting Information	Details
Date Reported	Date case was first reported to jurisdiction (mm/dd/yyyy).
Reporting Jurisdiction	State, territory, or jurisdiction reporting case to CDC.
State Case ID	Unique identifier given by the state health department.
Reporter Name, Phone Number, and Email	Contact information for person reporting case to CDC.
Clinician Name and Phone Number	Primary health care provider name and phone number.
Patient Status	Recurrent melioidosis is defined as a re-presentation with <i>B. pseudomallei</i> culture-positive clinical disease occurring <18 months following initial diagnosis and after the time designated for treatment completion (both intravenous and oral phases) for the previous episode, irrespective of whether the patient was adherent to the therapy or initially lost to follow-up.
Pathogen	Specify Burkholderia species.
Outbreak?	Denote if this case is part of a cluster or outbreak.

Case Demographic Information	Details
Sex	Genetic sex of patient.
Pregnant	Pregnancy status at onset of current illness.
Age	Age of patient at onset of current illness.
Residence	State, county, and zip code of patient's current residence.
Country of Usual Residence	If patient is not a US resident, denote country where patient usually resides.
Country of Birth	Indicate original country of birth, including US born. If unknown, please enter "Unknown."
Time in US	If not US born, indicate number of years patient has lived in the US.
Race and Ethnicity	Race and ethnicity of patient as noted in the chart or reported by physician or infection control personnel (ICP). Multiple boxes for race may be checked. Do not make assumptions based on name or native language. If race or ethnicity is unknown, please select "Unknown."
Occupation	Indicate occupation at time of disease onset. Specify past occupation(s) if relevant (i.e., occupations with environmental, animal, or travel related exposures).

Case Medical History	Details				
Pre-existing Medical Conditions Select all pre-existing medical conditions. If patient has no underlying medical conditions, select "No pre-existing conditions." Excessive alcohol use includes binge drinking (4+ drinks on an occasion for a woman or 5+ drinks on a					
Excessive Alcohol Use	Excessive alcohol use includes binge drinking (4+ drinks on an occasion for a woman or 5+ drinks on an occasion for a man) or heavy drinking (8+ drinks per week for a woman or 15+ drinks per week for a man).				



U.S. Department of Health and Human Services Centers for Disease Control and Prevention

Case Exposure Information	Details
Travel	Indicate all continents and US gulf coast states the patient has visited or lived in during their lifetime. Provide travel specifics for any travel in the 30 days prior to onset of current illness, if applicable.
Environmental Exposures	Indicate any water, mud, soil, compost, or sewage contact the patient had in the 30 days prior to onset of current illness and the locations where this contact occurred.
Animal Contact	Indicate any animal contact the patient had in the 30 days prior to onset of current illness, the type of animal, and the type of exposure.
Significant Weather	Indicate any significant weather events (e.g., monsoon, typhoon, cyclone, hurricane, flooding) experienced by the patient in the 30 days prior to onset of current illness.
Other Exposures	Specify any additional exposure information not captured elsewhere.

Case Clinical and Treatment Information	Details
Illness Onset	Date of the beginning of this illness (mm/dd/yyyy). Reported date of the onset of symptoms of this illness being reported to the public health system.
Symptoms and Conditions	Select patient-described symptoms or medically-identified conditions associated with this illness.
Hospitalization	Indicate whether the patient was admitted to a hospital for this illness. Enter admission and discharge dates, if applicable.
Treatment	Select the prescribed antimicrobial agents and duration for each. If prescribed other antibiotics, enter the generic name and duration, if known.
Post-Exposure Prophylaxis (PEP)	Indicate if the patient took PEP or the reasons for not taking PEP. If the patient took PEP, indicate if the patient completed the entire course of PEP as prescribed.
Outcome	Indicate the outcome of the patient following this illness. If the patient died of this illness, enter date of death.

Laboratory Testing Information*	Details
Test Type	Indicate the laboratory test performed.
Performing Laboratory	Indicate the laboratory that performed the test.
Specimen Type	Indicate the type of specimen collected.
Specimen Collection Date	Indicate the date the specimen was collected (mm/dd/yyyy).
Results	Indicate if the test was positive, any applicable qualitative results associated with the test, the species identified if applicable, and the test result date (mm/dd/yyyy).
Specimens to CDC	Indicate if the specimen was sent to CDC for testing.
AST Request	Indicate if the jurisdiction would like CDC to perform antimicrobial susceptibility testing on this specimen or isolate.

***NOTE:** Complete a new test block (4 available on the form) for each test performed.

Case Classification and Comments	Details
Case Classification	Indicate the patient's case classification based on the melioidosis case definition. Confirmed and Probable melioidosis cases must be reported to CDC following the notification criteria outlined in the CSTE position statement (22-ID-08).
Comments	List any other pertinent information about the case not provided elsewhere on the form.



BURKHOLDERIA SPP. INFECTION CASE INVESTIGATION FORM

Form Version Apr 2023

		NFO		

Date Reported: Reporti	ing Jurisdiction:	State C	ase ID:
Reporter Name:	Reporter Phone Nu	mber: Reporter Emai	:
Clinician Name:		Clinic	ian Phone Number:
		Other:	Part of an outbreak? Yes No Unknown
Save Mala Famala			Years Months Davs
Sex: Male Female		Age:	
Pregnant: Yes No Unknown		County:	-
Country of Usual Residence:	Co	untry of Birth:	Years in US:
Asian Na	ack or African American ative Hawaiian or Pacific Islander nknown	Other:	Ethnicity: Hispanic Non-Hispanic
Occupation:		r:	Unknown
Does the patient have any of the following p Diabetes Liver disease Malignancy Thalassemia On immunosurpressive drugs:	MEDICAL pre-existing medical conditions? Chronic lung disease Systemic lupus erythematosus	(select all that apply) Chronic kidney disease	No pre-existing conditions e Unknown
Does the patient excessively use alcohol or Current excessive alcohol use No Former excessive alcohol use Ur	o nknown		
	TRAVEL	HISTORY	
Has the patient EVER traveled or lived outs	side of the US in the lifetime (inclu	iding military service)? Yes N	lo Unknown
If yes, select all continents where patient has visited or lived in their	Asia Year:	Europe Year: Nort	h America (outside US) Year:
lifetime and most recent year visited:			ral America Year:
	Australia Year:	Caribbean Year: Sout	h America Year:
Has the patient served overseas in the milit	•		
Has the patient EVER visited or lived in any Alabama Florida Louisiana	of the following US states in the Mississippi Texas	ir lifetime? No/None Unknown	Year most recently visited:
In the <u>30 days prior to illness onset</u> , did the	e patient travel 50 miles or more fi	rom their normal residence? Yes	No Unknown
If yes, where?		Dates of Travel:	to:
If yes, where?			
If yes, where?		Dates of Travel:	
		O ANIMAL EXPOSURES	
In the <u>30 days prior to illness onset,</u> did the			Yes No Unknown
If yes, select all that apply: Running water (e.g., river, stream) Rainwater run-off/puddles	Still water <i>(e.g. lake, pond)</i> Mud or wet soil		eavy rainfall Sewage her soil
Specify locations where contact occurred:			
1	2	3	

Public reporting burden of this collection of information is estimated to average 10 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC/ATSDR Information Collection Review Office, 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30329-4027; ATTN: PRA (0920-0728).

In the <u>30 days prior to illness</u>	onset, did the pati	ient own or l	nave contact w	ith any animals'	? Yes	No U	nknown
If yes, select all that apply: Iguana Fish	Cat	Deg	Goat	Othory			
lguana Fish Sheep Horse	Mule	Dog Cow	Pig	Other			
Type of exposure:							
Handling or petting	•	esent in hom	e/property but	never touch	Location of p	ourchase or v	vhere animal was acquired:
Contact with animal fluids Cleaning enclosure/beddir	Other						
What activities led to the indicated environmental or animal exposure(s)? [select all that apply]	Swimming or Fresh water fi Adventure rac or mud run Biking/motoro Pet or livestoo	shing ce, triathalon cycle riding ck ownership	, Pla , Ga Pet farr o Dri	mping or hiking ying sports in ya rdening or yard v tting/touching ar n/zoo/other loca nking water	work nimals at	Washing Occupatio Other:	
	Boating, kaya	king, or rafti	ng Hu	nting		Unknown	
In the 30 days prior to illness	onset, has the pat	ient been in	any areas exp	eriencing signifi	cant weather?	Yes	No Unknown
If yes, select all that apply: Hurricane, cyclone, or typl Mudslide		looding/heav arthquake	vy rain		Vindstorm or tor)ther:	rnado	
Specify location:							
Please list any additional exp	osure information	not capture	d above:				
			INFORMATI	ON AND PR	ESENTATION	N	
Date of Illness Onset:							
Select all symptoms and con	ditions experience	d by the pat	ient during this	illness:			
Fever		nonia/pleura			ary infection		Ulcer
Nodule Anorexia		or soft tissue or joint infec		Septic sho Fatigue	ock		Respiratory distress Disorientation
Seizure	Joint p		lion	Chest pain			Weight loss
Periocardial effusion	•	abscess	c .	Headache			Sepsis
Muscle aches Skin abscess	Abdor	ninal discom	ifort	CNS infect	tion		Encephalomyelitis/meningitis/ extra-meningeal disease
Other symptoms or condit	ions:						-
		Т	REATMENT	AND OUTCO	DME		
Was the patient	Yes	1 st Admiss	sion Date:	1 st Di	scharge Date:		
hospitalized for	No	No 2 nd Admission Date:					
this illness?	Unknown	Unknown 3 rd Admission Date:		5			
		3 [.] Admiss	sion Date:	3 ^{.«} Di	scharge Date:		
Were antibiotics	Yes	Cefta	azidime		S	Start Date:	End date:
prescribed or administered to the patient?	No Unknown	Merc	openem		S	Start Date:	End date:
		Trim	ethoprim/Sulfar	nethoxazole	S	Start Date:	End date:
		Amo	xicillin/Clavular	nate	S	Start Date:	End date:
		Othe	er:		S	Start Date:	End date:
Did patient receive post-	Yes	If notion	t did not rocci	ve PEP, why not	2		
exposure prophylaxis	No	•	indicated	Allergic		er:	
(PEP)?	Unknown		ware of exposu	0			
		Unav	vailable	Unknov	wn		
If yes, antibiotic taken:			•	plete the cours	e? If patient	did not com	olete course, provide reason:
			∕es lo				
		(Jnknown				
Clinical outcome:	Died	(
Clinical outcome:	Died Still hospitaliz		Inknown Recovered Long-term	disability	Date of Death:		

Type: Specify tissue type: collection d Qualitative result: Positive Negative Borderline Indeterminate Other: Quanitative Negative Collection d Organism name: Lab result date:				LABO	RATORY TESTING I	NFORMATION				
HA ImmunoDot/DotBiot IgM Culture Other: Culture Performing lab: Specime Sp	1st Test & Sp	pecimen								
Specimen lype: Whole blood Serum Createrospinal fluid Serum Specify tissue type: Specify content to content	Test type:			1	Ũ		ated clinica	al laborat	ory system	
Speciment Vype: Whole blood Sarum Cerebrospinal fluid Tissue Specify tissue type: Specify contractive collection of collection of colle		Performing la	ab:							_
Qualitative Indeterminate Organisem name: Condentine Indeterminate Indeterminate Other: Quantitative result (a.g., titer): Send to CDC? Yes No, isolate destroyed No, specimen not available AST requested? Yes No Not applicable 2nd Test & Specimen Test type: PCR IHC Other ELISA IgM Viteck or other automated clinical laboratory system 9erforming lab: Performing lab: Specify tissue type: Specify tissue type: Specify clinical laboratory system 0ualitative result: Positive Borderline Indeterminate Other: Quantitative result (e.g., titer): Collection of collection of collection of transmate 0ualitative result: Positive Borderline Indeterminate Other: Quantitative result (e.g., titer): Collection of collection of collection of transmate 0rganisem name: Unine Other: Quantitative result (e.g., titer): Specify tissue type: Lab result date: Serving lab: PCR IHC Other ELISA IgM Viteck or other automated clinical laboratory system Test type: PCR IHC Other: Quantitative result (e.g., titer): Specime Specimen Whole blood Cerebrospinal fluid Carture<	•	Serum	Tissue			1 5 51			co	Specimen blection date:
Send to CDC? Yes No, isolate destroyed No, specimen not available AST requested? Yes No Not applicable 2nd Test & Specimen Test type: PCR IHC Other ELISA IgM Culture Viteck or other automated clinical laboratory system Other: Specimen Specimen Urine Viteck or other automated clinical laboratory system Culture Specimen Culture Specimen Urine Specimen Urine Specimen Collection of Culture Specimen Collection of Culture Specimen Urine Specimen Urine Specimen Urine Culture Culture Culture Culture Culture Culture Culture Specimen Collection of Culture Specimen Collection of Culture Culture No No No Applicable 3rd Test & Specimen type: PCR IHC Other ELISA IgM Culture Culture Viteck or other automated clinical laboratory system Culter: Collection d 4urine Other: Culture Culture Viteck or other automated clinical laboratory system Culter: Collection d 4urine Other: Culture			Borderline				Quantitat	ive result	: (e.g., titer):	
Send to CDC? Yes No, isolate destroyed No, specimen not available AST requested? Yes No Not applicable 2nd Test & Specimen Test type: PCR IHC Other ELISA IgM Culture Viteck or other automated clinical laboratory system Other: Specimen Specimen Urine Viteck or other automated clinical laboratory system Culture Specimen Culture Specimen Urine Specimen Urine Specimen Collection of Culture Specimen Collection of Culture Specimen Urine Specimen Urine Specimen Urine Culture Culture Culture Culture Culture Culture Culture Specimen Collection of Culture Specimen Collection of Culture Culture No No No Applicable 3rd Test & Specimen type: PCR IHC Other ELISA IgM Culture Culture Viteck or other automated clinical laboratory system Culter: Collection d 4urine Other: Culture Culture Viteck or other automated clinical laboratory system Culter: Collection d 4urine Other: Culture		Organism nar	me:					Lab	result date	
Test type: PCR IHA IHC ImmunoDot/DotBiot IgM Culture Other ELISA IgM Culture Vitack or other automated clinical laboratory system Other: Specimen type: Whole blood Cerebrospinal fluid Serum Specime Tissue Specime Collection of Collection	Send to CDC?	Yes N	lo, isolate destroyed	No, s	pecimen not available	AST requested?	Yes	No	Not app	licable
IHA ImmunoDot/DotBlot IgM Culture Other: Performing tab:	2nd Test & S	pecimen								
Specimen type: Whole blood Serum Cerebrospinal fluid Insue Specify tissue type: Specime collection of collection of colle	Test type:	IHA	ImmunoDot/DotBlot IgN		-		ated clinica	al laborat	ory system	
result: Negative Indeterminate Other: Quantitative result (e.g., titer): Organism name:	•	Whole bloc Serum	od Cerebrospina Tissue	l fluid					C(Specimen ollection date
Send to CDC? Yes No, isolate destroyed No, specimen not available AST requested? Yes No Not applicable 3rd Test & Specimen PCR IHC Other ELISA IgM Viteck or other automated clinical laboratory system Test type: PCR IHC Other ELISA IgM Viteck or other automated clinical laboratory system Specimen Whole blood Cerebrospinal fluid Specify tissue type: Specime collection d Qualitative result: Positive Borderline Specimen not available AST requested? Yes No Not applicable Gualitative result: Positive Borderline Specify tissue type: Quantitative result (e.g., titer):				Other: _			Quantitat	ive result	(e.g., titer)	·
3rd Test & Specimen Not reference of the opproduct of the oppr		Organism na	me:					Lab	result date	
Test type: PCR IHA IHC ImmunoDot/DotBlot IgM Other ELISA IgM Culture Viteck or other automated clinical laboratory system Other: Specimen type: Whole blood Serum Cerebrospinal fluid Tissue Specify tissue type: Specify tissue type: Specime collection d Qualitative result Positive Negative Borderline Indeterminate Other: Quantitative Culture Quantitative AST requested? Yes No Not applicable 4th Test & Specimen type: PCR IHA IHC Other: Other ELISA IgM Culture Viteck or other automated clinical laboratory system Culture Specify tissue type: Callection d Guantitative result Positive Negative Borderline Indeterminate Other: Quantitative Culture Specify tissue type: Callection d Specimen type: PCR IHA IHC Other ELISA IgM Culture Viteck or other automated clinical laboratory system Other: Specime Collection c Specimen type: Whole blood Cerebrospinal fluid Serum Specify tissue type: Specime collection c Quantitative result Positive Negative Borderline Indeterminate Other: Quantitative result (e.g., titer): Quantitative result (e.g., titer):	Send to CDC?	Yes N	lo, isolate destroyed	No, s	pecimen not available	AST requested?	Yes	No	Not app	licable
IHA ImmunoDot/DotBlot IgM Culture Other:	3rd Test & Sp	becimen								
Specimen type: Whole blood Serum Cerebrospinal fluid Tissue Specify tissue type: Specify content collection of collection	Test type:			1			ated clinica	al laborat	ory system	
Opecified White block Objecting in third Specify tissue type: collection d type: Serum Tissue		Performing la	ab:							
result: Negative Indeterminate Other: Quantitative result (e.g., titer): Organism name:	•	Serum	Tissue						cc	Specimen Ilection date:
Send to CDC? Yes No, isolate destroyed No, specimen not available AST requested? Yes No Not applicable 4th Test & Specimen PCR IHC Other ELISA IgM Viteck or other automated clinical laboratory system Test type: PCR IHC Other ELISA IgM Viteck or other automated clinical laboratory system Performing lab: Performing lab: Specimen Specime collection of the collection				Other: _			Quantitat	ive result	(e.g., titer)	
4th Test & Specimen PCR IHC Other ELISA IgM Viteck or other automated clinical laboratory system Test type: PCR IHC Other ELISA IgM Viteck or other automated clinical laboratory system Performing lab:		Organism na	me:					Lab	result date	
Test type: PCR IHC Other ELISA IgM Viteck or other automated clinical laboratory system IHA ImmunoDot/DotBlot IgM Culture Other: Other: Performing lab: Cerebrospinal fluid Specime Specify tissue type: Specime type: Vine Other: Other: Collection of the	Send to CDC?	Yes N	lo, isolate destroyed	No, s	pecimen not available	AST requested?	Yes	No	Not app	licable
IHA ImmunoDot/DotBlot IgM Culture Other: Performing lab:	-	1								
Specimen type: Whole blood Serum Cerebrospinal fluid Specify tissue type: Specify collection of collection	Test type:	IHA	ImmunoDot/DotBlot IgN	1			ated clinica	al laborat	ory system	
Qualitative result: Positive Negative Borderline Indeterminate Quantitative result (e.g., titer):	•	Whole bloc Serum	od Cerebrospina Tissue						C	Specimen
				Other:			Quantitat	ive result	(e.g., titer):	
Organism name: Lab result date:		Organism na	me:					Lab	result date	
Send to CDC? Yes No, isolate destroyed No, specimen not available AST requested? Yes No Not applicable	Send to CDC?	Yes N	lo, isolate destroyed	No, s	pecimen not available	AST requested?	Yes	No	Not app	licable
CASE CLASSIFICATION					CASE CLASSIFIC	ATION				
Confirmed Probable Suspect Not a case Unknown		Confirme	d Probab	le	Suspect	Not a case		Unknow	/n	
ADDITIONAL COMMENTS					ADDITIONAL COM	IMENTS				

DOB: date of birth, PCR: polymerase chain reaction, IHA: indirect hemagglutination, IHC: immunohistochemistry, AST: antimicrobial susceptibility testing