BRUCELLOSIS CASE REPORT FORM





Brucellosis Case Report Form General 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.

Send the completed form with all <u>personal identifiers removed</u> to:

Mail: Centers for Disease Control & Prevention

ATTN: Bacterial Special Pathogens Branch

Mailstop C09

1600 Clifton Rd NE Atlanta, GA 30333

Fax: (404) 639-7080

Patient identifier information (NOT transmitted to CDC)

Patient Name	Patient's full name
Phone	Patient's phone number
Patient Chart Number	Medical chart number for patient
Address	Patient's address including street and city
State, Zip	Patient's state of residence and zip code
Hospital Name	Name of the hospital where the patient is admitted or seen

Information obtained for confirmed and probable brucellosis cases

PATIENT & PHYSICIAN INFORMATION

TATIENT & THISICIEN INFORMATION							
State Case ID	Unique identifier given by the state health department.						
Investigator	State health department investigator name.						
Date Reported	Date the case was reported to state.						
Physician	Primary health care provider name.						
Phone	Primary health care provider phone number and/or pager.						
NETSS Number	If case submitted to NETSS, include the NETSS-generated Case ID number.						

DEMOGRAPHICS

State of Residence	Use the 2 letter postal abbreviation (e.g., NY) of patient's state of residence.
County of Residence	Patient's county of residence.
Age	Age of patient at time of diagnosis; indicate age unit as months or years.
Sex	Genetic sex of patient (i.e., male or female).
Pregnant	Pregnancy status at time of diagnosis.
Country of Birth	Indicate original country of birth, including U.S. born. If unknown, please enter "Unknown".
Ethnicity	Indicate ethnicity of patient.
	Race of patient as noted in the chart or reported by physician or infection control personnel (ICP). Multiple
Race	boxes may be checked. Do not make assumptions based on name or native language. If race is unknown,
	please check "Unknown".
Occupation	Indicate occupation at time of disease onset. Specify past occupation(s) if relevant.

CLINICAL INFORMATION AND TREATMENT

Disease Presentation	Disease presentation- a date determined by duration from onset of symptoms to date of diagnosis.
Symptoms and	Select patient-described symptoms. Enter date of onset if known. If approximate date is known, enter
Associated Diagnoses	rounded date (e.g., fever two weeks prior to seeking medical care on 9/17—enter 9/1).
Signs and	Select signs identified upon examination. Enter date of diagnosis where known. Enter an approximate date if
Associated Diagnoses	a precise date is unknown.
Hospitalized?	Indicate whether the patient was admitted to a hospital due to this illness. Enter admission and discharge date,

	if applicable.
Deceased?	Indicate if the patient died of this illness. Enter date if applicable.
Treatment and Duration	Select whether the patient has completed their treatment. Select the prescribed antimicrobial agents, amount, and duration for each. If prescribed other antimicrobials, enter the generic name, amount, and duration, if known. NOTE: If an agent is taken twice daily, enter the total prescribed mg/day (e.g., 100 mg BID- enter 200 mg/day).

RISK FACTORS

Travel	Select whether the patient traveled out of state or country in the past six months, and where and when if applicable.
Animal Contact	Select which animals and type of contact, if any, the patient had in the past 6 months.
Unpasteurized Dairy	Select if the patient consumed unpasteurized (raw) dairy in the past six months. Choose type of animal, owner of the animal the dairy came from, what products were eaten, and location of product.
Confirmed Case	Select if the patient is linked to a confirmed case. If yes, select the relationship to the patient.
Similar Illness	Select if the patient is aware of a contact having a similar illness. If yes, select the relationship to the patient.
Risk Status	If the patient had a known exposure to <i>Brucella</i> , indicate the exposure source and the location of exposure. Also indicate the assessed risk status of the exposure. Finally, if exposed to a <i>Brucella</i> vaccine, indicate to which vaccine the case was exposed. The CDC exposure guidelines are available at www.cdc.gov/mmwr/preview/mmwrhtml/mm5702a3.htm . If a laboratory exposure did occur, review these assessment, monitoring, and prophylaxis recommendations. For assistance, contact CDC at the phone number listed on page one.
Received Post-Exposure Prophylaxis (PEP)	If the patient was exposed to <i>Brucella</i> , indicate if the patient took PEP, or reasons for not taking PEP.
Completed PEP	If exposed, indicate if the patient completed the entire course of PEP as prescribed. CDC recommended PEP regimen is doxycycline 100 mg orally twice a day plus rifampin 600 mg orally once a day for 21 days.

LABORATORY DATA

w Laboratory Data section for each laboratory receiving and processing patient samples.								
field blank for each test not performed.								
Indicate case classification. Confirmed and Probable cases must be reported to NETSS by the next regularly scheduled transmission cycle. CDC must be notified of multiple cases which are temporal/spatial clusters within 24 hours of the cases meeting the notification criteria (CSTE Position Statement 09-SI-04).								
Name Enter the laboratory name and address which processed the sample. For each laboratory that processed the sample, start a new laboratory section. Submit a copy of page four for each laboratory involved in testing.								
Enter the name, city, and state of the laboratory from which the specimen is received; include date of receipt.								
If a paired agglutination test was done, enter results in this table. If known, enter the agglutination test (SAT, BMAT, Tube AT). Indicate which titers were run- total antibody (complete) and/or IgG (reduced). Enter in the acute and convalescent titers. Indicate if one, both, or paired titers are positive. Enter the testing laboratory's positive cut-off value for the test. If a single titer was done, enter as an acute titer. For ELISA, indicate if IgG, IgM, or both titers were run. Enter in the acute and convalescent titers and if one, both, or paired titers are positive. Enter the testing laboratory's positive cut-off value for the test.								
Enter the dates the acute and convalescent samples were collected.								
Enter the value or titer in the row of the test completed, and whether the test was considered positive. If the test used is not listed, enter name and results in "Other". Indicate the laboratory's positive cut-off value for the test.								
Select whether PCR and/or culture was attempted. Indicate the source of specimen used for the specified test. Enter the date of specimen collection, if the test was positive, and the species identified (e.g.: <i>abortus</i> , <i>canis</i> , <i>melitensis</i> , <i>suis</i> , other).								
Indicate if the specimen for culture was collected prior to administration of antimicrobial therapy.								
Indicate if a culture-positive result of a select agent was reported to CDC, as required by regulation. Reporting requirements and forms are available at http://www.selectagents.gov/ .								
Select if laboratory workers were possibly exposed during specimen processing. The CDC exposure guidelines are available at www.cdc.gov/mmwr/preview/mmwrhtml/mm5702a3.htm . If a laboratory exposure did occur, review these assessment, monitoring, and prophylaxis recommendations. For assistance, contact CDC at the phone number listed on page one.								
If a laboratory exposure occurred, indicate if the "release" of a select agent was reported to CDC, as required by								
regulation. Reporting requirements and forms are available at http://www.selectagents.gov/ .								
Indicate if the specimen was sent to CDC for testing.								
Indicate if the specimen is still available, if needed for future testing.								

Case Name	Phone	Medical Chart No	



DEPARTMENT OF HEALTH AND HUMAN SERVICES Centers for Disease Control and Prevention Atlanta 6,3 3033 Remove case identifier information prior to transmission to CDC.

BRUCELLOSIS CASE REPORT FORM

Form Approved
OMB No. 0920-0004
Exp. Date 6/30/2013



Atlanta, GA 30033												Ехр.	Date 6/30/2013	CENTERS FOR DISEASE' CONTROL AND PREVENTION
			- (CASE	& PHYS	SICIAN	INFO	PRMATION	-					
State Case ID	Physician						Phone Number							
Investigator			NETSS	ID No	(if notifie	ed):	 SE ID	·——— -	SITE	 ST	ATE			
Date Reported					- DEM	IOGRA			SITE	31	AIL			
State of Booklanes	Course	tu of Do	aidaa						D.		Cov. 🗆	Mala		l Halmanina
State of Residence														
Pregnant Yes No Unknown Country of Birth Ethnicity Hispanic Non-Hispanic Unknown														
Race ☐ American Indian/ Alaskan Native Occupation ☐ Animal research ☐ Medical research ☐ Dairy ☐ Laboratory ☐ Wildlife ☐ Asian/Pacific Islander ☐ Black ☐ White ☐ Rancher ☐ Slaughterhouse ☐ Tannery/rendering ☐ Veterinarian/Vet Tech ☐ Lives with person of above occupation ☐ Other														
- CLINICAL INFORMATION AND TREATMENT -														
Disease Presentation ☐ Acute (0-8 weeks) ☐ Subacute (8 weeks - <1 yr) ☐ Chronic (1 yr+) ☐ Unknown														
Symptoms, Signs, and A	ssociate	ed Diagr	noses (in	dicate	date of	onset	or di	agnosis):						
Yes No Unk Symptom	Date Ons	et	Yes No	Unk Sy	ymptom/	Sign	Date	Diagnosis	Yes N	o Unk	Signs		Date of	of Diagnosis
□ □ □ Fever	/_/	/		□ Ai	norexia		/	/			Hepato	megaly		/
Max temp:				_	yalgia			/				megaly		
□ □ □ Night sweats				_				/			Arthritis			
1			<u> </u>	_ ''	eight lo									/
	//				ndocard	itis		/			Mening		/_	'
	//	/			rchitis		/-	/			Spond		/_	/
□ □ □ Fatigue	//	<u>' </u>		<u> </u>	pididym		/_	/		J U	Other:_			/
Was the case hospitalized	П,	Vac \square N	lo □ Unk	nown		ŀ	f yes,	admission da	ate:			/-	/	
because of this illness?		163 🗆 1	NO LI OTT	TIOWIT		1	f appli	cable, discha	arge dat	e:		/	/	
Is the case deceased?		Yes □ N	lo □ Unk	nown		1	f yes,	date of death	h:			1	/	
Treatment and Duration (ch	neck all th	at apply): 🗆 Cur	rently u	ınder tre	atment		Completed tr	eatment		Not treate			
□ Doxycyclinemg				☐ Oth									ng/day	days
☐ Rifampinmg				□ Oth									ng/day	days
☐ Streptomycin mg	J/day	days		□ Oth	er:							r	ng/day	days
	_	-			- DIS	SK FAC	TOP.						_	
In the 6 months prior to	illness	onset	did the	case		K FAC	TOR	S -						
Travel outside state of residence			•											
If Yes, where?				01		D	ates o	of travel	1 1	/	to	1 1		
If Yes, where?						D	ates o	of travel	1		to	11		
Have contact with animals?	<u>?</u> □ Yes	□ No □	□ Unknov	vn					Wh	o ow	ns the a	nimal(s)?	
Type of contact			at Shee					Other		ase	Private	Wild	Commercia	
Birthing/animal products									_					
Skinning/slaughter									_					
Hunting									⊣ ⊢					
Other:									_L_L					
Consume unpasteurized d					Yes □								e product ac	quired?
Type of food product		Pig Go			_		+	Other		.S.		her		Other
Milk				\perp				<u> </u>						
Fresh/soft cheese								<u> </u>						
Undercooked meat							 	<u> </u>						
Other:														
Have a link to a confirmed case?												ıa ⊔N	ieignbor ⊔ (oworker
			C NI~	Know of similar illness in contact?										
Know of similar illness in o	contact?				nown	144	au'.		linical a-			rator: '	7 Form/Don	
Know of similar illness in of Have an exposure	contact? Clinical sp	ecimen	☐ Isolat							etting	□ Labo	-		:h
Know of similar illness in of Have an exposure to a Brucella?	contact? Clinical sp /accine	ecimen □ Unkı	☐ Isolat nown	e		expos	ure o	ccur? 🗆 St	urgery	etting Un	 □ Labo known	☐ Othe	r:	
Know of similar illness in of Have an exposure to a Brucella?	contact? Clinical sp /accine	ecimen Unki	☐ Isolat	e	If ex	expos	ure oo to va	ccur?	urgery ate which	etting Un	□ Labo known □ S19	□ Othe		□ Other

If yes, did ca	se complete co	urse? □ Yes □	No □ Unknov	wn E] Partial <i>explai</i>	n:					
Public reporting burden of this collection of information is estimated to average 20 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 30333; ATTN: PRA (0920-0004).											
-CASE DEFINITION (2010) -											
Confirmed: A clinically compatible illness with definitive laboratory evidence (i.e.: culture and identification of <i>Brucella</i> spp. from clinical specimens OR serological evidence of a fourfold rise in <i>Brucella</i> antibody titer in paired acute and convalescent serum specimens greater than or equal to 2 weeks apart). Probable: A clinically compatible illness epidemiologically linked to a documented <i>Brucella</i> case OR has presumptive laboratory evidence (i.e.: <i>Brucella</i> total antibody titer of greater than or equal to 160 by standard tube agglutination test (SAT) or <i>Brucella</i> microagglutination test (BMAT) in one or more serum specimens obtained after onset of symptoms OR detection of <i>Brucella</i> DNA in a clinical specimen by PCR assay). - LABORATORY DATA -											
Leave	the test field bla	nk for each test not	performed.			orocessir	ng case samples.	Print e	xtra copies if necessary.		
		rmed ☐ Serologio									
Laboratory Na	me:		_ City:				State:		Zip:		
Received Fron	າ:		_ City:			State:_	Date	Recei	ved://		
								he test	t was not performed.		
Paired Ser	ologic Tests	Titers	Acute Tite	r	Convalesce	nt Titer	Positive	?	Positive Cut-off:		
Agglutinatior Test:	1	☐ Total antibody ☐ IgG				_ ☐ Yes ☐ No ☐ Unkn ☐ Yes ☐ No ☐ Unkn					
ELISA:		□ IgG □ IgM	_:				☐ Yes ☐ No ☐ Unknown ☐ Yes ☐ No ☐ Unknown				
Date Sample	Collected:	Acute://	Con	vales	cent:/	/					
Other Se	rologic Tests	Titer or Value	Po	ositiv	e?	Pos	sitive Cut-off				
Rose Bengal		_:	□ Yes □	No	□ Unknown						
Coombs IgG		_:	☐ Yes ☐ No ☐ Unknown								
Other:		:	□ Yes □	No	o □ Unknown						
Other:		()	□ Yes □	No	□ Unknown						
Other Tests	s	ource of Specime		Dat	e Collected		Positive?		Species		
PCR □ Blood □ Abscess/wound □ Bone Marrow □ CSF □ Other:					//		☐ Yes ☐ No ☐ Unknow				
Culture Blood Abscess/wound Bone Marrow/ Pes No Unknow						vn					
Was the spec	cimen for culture	collected prior to a	antimicrobial the	erapy	?	☐ Yes	s □ No □ Unknov	vn			
If culture positive, was the identification of a select agent reported to CDC? ☐ Yes ☐ No ☐ Unknown											
Did a possibl	e laboratory exp	osure occur?	Yes □ No □	Unkno	own If ye	s, was it	reported to CDC?		Yes □ No □ Unknown		
	Were specimens sent to CDC for testing? ☐ Yes ☐ No ☐ Unknown Is the specimen still available? ☐ Yes ☐ No ☐ Unknown										