Attachment E. Screen shots for Burden of Canine Brucellosis Information Collection Instrument

Landing page (page 1):



Form Approved OMB No. xxxx-xxx Expires xxxxxxxx

Burden of Canine Brucellosis Information Collection

Welcome! Thanks for taking the time to participate in this information collection. Below you will find a brief description of the project. Please review this description before proceeding through the information collection instrument.

Project Title: Burden of Canine Brucellosis

Investigators: Rita Traxler, Dr. Marta Guerra, Centers for Disease Control and Prevention

Protocol and Consent Title: Burden of Canine Brucellosis Information Collection

You are being asked to volunteer in an information collection. This information collection will ask about Brucella canis lab results from dogs. It will also identify lab tests used, lab exposure policies, and state reporting policies. Average time for completion should not exceed one hour. Information collected through this process will be kept secure, and will be presented in aggregate form, so as not to include any individually identifiable information.

Purpose: To assist in estimating the burden of canine brucellosis in the United States. **Requirements:** You must be age 18 years or older to participate in this information collection

Procedures: Some of the information collection questions will require review of your records. To save time we recommend printing a copy of the PDF information collection insrument attached to the email you received, then entering the data into the online collection instrument. Please click on "Next" to participate in the online information collection instrument. There are 22 questions. You will need to look up data in your results database. Please click "Submit" when you reach the end of the information collection instrument. If your lab does not conduct any testing on dogs, please start the information collection instrument and complete the first 4 questions, then submit the information collection instrument on the final page.

Confidentiality: No identifying information will be gathered. All responses will remain secure between the respondent and the project administrators. Responses will be aggregated in future publications.

Participant Rights: Your participation in this information collection is voluntary. You do not have to complete the information collection instrument if you don't want to. You have the right to change your mind and exit the information collection instrument at any time without giving any reason and without penalty.

Questions about the Information collection instrument: If you have any questions about the information collection instrument, you may contact Rita Traxler (rtraxler@cdc.gov) at (404) 639-0265.

Participant Agreement: By participating in the information collection instrument, it means that you have read (or have had read to you) the information given in this consent form, and you would like to be a volunteer in this information collection.

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	atory Nar	ne	
	•		
2. Labora	tory Stat	e	
3. Descri	be your la	aboratory (check all tha	t apply):
	State univ	versity-based laboratory	
	_	ernment facility (e.g., Dep	t. of Agriculture lab)
_	Federal fa	,	
	-	Animal Health Laboratory N	etwork lab
	Other, Sp	ecify:	
Yes	⊚ No	□ Unknown	
		n electronic database	for specimen information and data
5. Do yo storage	?		
	?	Unknown	
storage Yes 6. If yes	No No s, how ma		nother compatible electronic syste
storage Yes 6. If yes	No No s, how maplace in	any years has this or a	nother compatible electronic syste
Storage Yes 6. If yes been in < 1 year	No No ma	any years has this or a your laboratory?	

Public reporting burden of this collection of information is estimated to average 1 hour 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-xxxxx).

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		· · · · · · · · · · · · · · · ·		your lab during t	the last:	years
Jan 1, 20)10 - Dec 3	1, 2014; or	if <5 years, use	this date range		
	a. Did you	outsource	testing for any o	f these samples?		
	○ Yes	○ No	Ounknown			
				id you outsource?		
	O 1-25%	○ 26-509	% ○ 51-75 %	○ 76-100%		
		h lab(s) did ate of each		esting during this	time frame? Lis	st the nam
	Lab Na	me	City		State	
					AK-Alaska	~ ::
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Jan 1, 2010 - Dec 31, 201	.4 or
a. What types of specimens are sub (give the number of each sample ty	omitted to your laboratory requesting culture? rpe, if available)
Blood	Fluid specify
Vaginal Swab	Other specify
Semen	Other specify
Tissue	Unknown Type
Urine	■ Unknown number of samples
c. now many dogs have you rep	orted out as culture positive for Brucella canis:
d. Have any dogs been culture p	positive for other Brucella species?

Da	90	5	
га	ςc.	J	•

Transfer to another	lab Destroy/discard	d	
Retain in inventory	Other		
	tains isolates in you isolates for assay d	ur inventory, would you consider allowing C evelopment?	DC
⊚ Yes		⊚ No	
Maybe - please	contact us to discuss	Unknown	
Of the canine	samples proces	ssed in your lab during the last :	yea
			_
Jan 1.	2010 - Dec 31, 201	L4 or -	
,	2010 - Dec 31, 201		
a. What types	of specimens are	submitted to your laboratory requesting PC	R for
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a. What types of specimens were submitted to your laboratory requesting serology for bacterial agents? (give the number of each sample type, if available) Serum	or
bacterial agents? (give the number of each sample type, if available) Serum CSF Unknown type Unknown number of samples i. Does this include paired samples? Yes No Unknown b. How many samples were sero-positive for rough Brucella species (i.e. B. canis)? Unknown c. How many dogs were considered sero-positive for Brucella species (e.g., Brucella suis)? Yes No Unknown d. Have any dogs been sero-positive for smooth Brucella species (e.g., Brucella suis)? Yes No Unknown No. of Dogs 12. If yes, how many samples, and how many dogs were positive? No. of Samples No. of Dogs 12. If your lab obtains a positive serology result, what do you do with the sera? Transfer to another lab Destroy/discard Retain in inventory Other a. If your lab retains serum in your inventory, would you consider allowing CDC access to sera for assay development? Yes No Waybe-please call to discuss Inknown Public reporting burden of this collection of information is estimated to average 1 hour per response, including the I reviewing instructions, searching costing data sources, gathering and maintaining the data needed, and completing in reviewing instructions, searching costing data sources, gathering and maintaining the data needed, and completing in reviewing instructions, searching costing data sources, gathering and maintaining the data needed, and completing in reviewing the cultication of information is estimated to average 1 hour per response, including the I reviewing instructions, searching confirmation contacts of information in a searching confirmation of the required to report to reviewing the cultication of information is estimated to average 1 hour per response, including the I reviewing the cultication of information is estimated to average 1 hour per response, including the I reviewing the cultication of information is estimated to average 1 hour per response, including the II reviewing the cultication of the contact is averaged to the contact in some and the cultication of the contact is averaged to the com	or
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7: Agar Gel Immunodiffusion Test (AGID) Screening Confirmatory Experimental Diagnost Indirect Fluorescent Antibody Test Screening Confirmatory Experimental Diagnost (IFAT or IFA)	
ELISA Screening Confrimatory Experimental Diagnost	tic
PCR Screening Confirmatory Experimental Diagnost	

Screening Confirmatory Experimental Diagnostic

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Other, specify

15. If you have more details that you would like to share about the laboratory results section, please comment here $\,$

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	a. In do	gs?					
	Yes	⊚ No	Unkn	own			
	i. If	yes, to who	m do you	report a posit	tive result?	,	
		State	Dept. of	Agriculture			
		State	Bureau o	of Animal Hea	alth		
		US D	ept. of Ag	riculture			
		Othe	r <i>please</i>	specify			
	b. In h	umans?					
		No	Unk				
		e number (of lab wo	orkers expo			n your lab
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YesNo	Not written but do have an informal plan
a. [If yes or Informal plan] Se informal policy:	elect the components of your written o
Symptom monitoring	■ Serological monitoring
■ Prophylactic antibiotics	■ Risk evaluation/classification
Consult occupational health	☐ Other specify
19. Does your lab have a writter canis exposures? Ores No	n policy specific to human B. Not written but do have an informal plan
canis exposures? Yes No	Not written but do have an informal plan
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	= Local Hashb Danashmant
	Local Health Department
	State Health Department
	Occupational Health/Infection Control Officer
	Centers for Disease Control and Prevention
	US Dept. of Agriculture
	Other:
Yes	⊚ No
Maybe ple	ase contact us to discuss Unknown
22. If you	have more details that you would like to share about the policy and section, please comment here:
22. If you	have more details that you would like to share about the policy and