Column E Explanation

This form is intended as an aid to complete the Column E explanation. It is not an official form and its use is voluntary. Annual Reports and explanations should NOT include PII information such as names (principle investigators and research staff), addresses, protocols, meeting notes (either in part of in full), the animals room numbers, grant information, veterinary care programs, and the like. A Column E explanation must be written so as to be understood by lay person as well as scientists.

1.	Registration Number: 31-R0028			
2.	Number of animals used in this study: 24			
3.	Species (common name): <u>Ferrets</u> of animals used in this study.			
4. Explain the procedure producing pain and/or distress. Explanations should include a bit description of the procedure, but also explain what the animal's experience, examples include, but are not limited to: Neurological signs, seizures, tremors, paralysis, lethargy inappetance, respiratory signs, GI distress, vomiting, and diarrhea.				
	This is an influenza disease model. Animals were infected intranasally with influenza virus. The appropriate inoculum was identified in Part 1 of the experiment. Part 2 of the experiment involves inoculation of animals and placement in 1 of 6 treatment groups. Upon successful inoculation ferrets presented with fever, nasal discharge, sneezing, coughing, and malaise. No dehydration was noted. Overall fevers were mild; beginning 2 days post infection temperatures ranged from $36.2-41^{\circ}\text{C}$, with an average of 38.2°C .			
5.	Attach or include with the reason(s) for why anesthetics, analgesics and tranquillizers could not be used. (For federally mandated testing, see Item 6 below).			
	This is a disease model where anti-infective drugs are being evaluated for efficacy. Anesthetics, analgesics, and tranquillizers are not warranted and may even worsen symptoms or lesson the effects of the treatments. Also, clinical signs of illness including temperature and activity level were assessed daily and used as indicators of treatment efficacy.			
6.	What, if any, federal regulation require this procedure? Cite the agency, the code of Federal Regulations (CFR) title number and the specific section number (e.g. APHIS, 9 CFR 113.102):			
	Agency N/ACFR_N/A			

2017 AR Attachments Submitter Notice Page 4 of 4

То:	2017 AR Review Coordinator	Facility Contact:	ami Mc Cour	
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Facility Fax:	(216) 368-4805	No. of Pages:	>	
Facility Phon	ne:(216) 368-4972	Date: Novemb	per 1, 201	
NO ANNUAL	REPORT ATTACHMENTS SUBMITT	ED TO AC FOR 2017		
☐ We did not	t submit AR Attachments to AC for 2017.			
NO OBJECT	IONS RESPONSE			
We have r	no objections to the release of our AR Atta	chments as received ar	nd do not intend to	
seek judicial re	eview to bar release of these documents.			
REDACTIONS	S PURSUANT TO EXEMPION 4 REQUE	STED		
☐ We object	to the release of our AR Attachments as r	eceived and ask that yo	ou consider the	
enclosed justification statement and suggested redactions.				
COMMENTS				
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	Me (I)	9	11/1/2017	
Signature		Date		
Mark	Chance, IO			
Print Name (i	if different from above)			