According to the Paperwork Reduction Act of 1995, an agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless According to the Paperwork Neducation According to 1995, an agency may not conduct or sponsor, and a person is not required to complete this information collection is 0579-0036. The time required to complete this information collection is estimated to average 2 hours per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data OMB APPROVED needed, and completing and reviewing the collection of information. This report is required by law (7 U.S.C. 2143). Failure to report according to the regulations can result in an order to cease and desist and to be subject to penalties as provided for in Section 2150. Interagency Report Control Fiscal Year, 2009 No. 0180-DOA-AN UNITED STATES DEPARTMENT OF AGRICULTURE REGISTRATION NUMBER: 33-V-0004 ANIMAL AND PLANT HEALTH INSPECTION SERVICE 2. HEADQUARTERS RESEARCH FACILITY (Name and Address, as registered with USDA ANNUAL REPORT OF RESEARCH FACILITY Edward Hines Jr Va Hospital (151) (b)(2)High, (b)(7)f Hines, IL 60141 NOV 2 7 2009 (TYPE OR PRINT) Telephone: (708) 216 2111 3. REPORTING FACILITY (List all locations where animals were housed or used in actual research, testing, teaching, or experimentation, or held for these purposes. Attach additional sheets if FACILITY LOCATIONS (Siles) See Attached Listing REPORT OF ANIMALS USED BY OR UNDER CONTROL OF RESEARCH FACILITY (Attach additional sheets if necessary or use APHIS FORM 7023A.) Number of animals upon Number of animals upon which teaching which experiments, experiments, research, surgery, or tests were Number of animals Number of animals teaching, research, conducted involving accompanying pain or distress to the animals and for which the use of being bred, surgery, or tests were conducted involving teaching, research, experiments, or Animals Covered By conditioned, or held appropriate anesthetic, analgesic, or tranquilizing drugs would have adversely affected the procedures, results, or TOTAL NUMBER OF ANIMALS The Animal for use in teaching. accompanying pain or distress to the animals and for which Welfare Regulations tests were testing, experiments. conducted involving research, or surgery but not yet used for interpretation of the teaching, research, experiments, surgery, or tests. (An explanation (Cols. C + D + E) no pain, distress, or appropriate and use of pain-relieving such purposes. analgesic, or of the procedures producing pain or distress on these animals and the reasons such drugs drugs. tranquilizing drugs were used. were not used must be attached to this report. 0 4. Dogs 5. Cats 6. Guinea Pigs 85 a 7. Hamsters 8 20 6 8. Rabbits 9. Non-human Primates 10. Sheep 0 11. Pigs 12. Other Farm Animals 13. Other Animals ASSURANCE STATEMENTS Professionally acceptable standards governing the care, treatment, and use of animals, including appropriate use of anesthetic, analgesic, and tranquilizing drugs, prior to, during, and following actual research, teaching, testing, surgery, or experimentation were followed by this research facility 2) Each principal investigator has considered alternatives to painful procedures. This facility is adhering to the standards and regulations under the Act, and it has required that exceptions to the standards and regulations be specified and explained by the principal investigator and approved by the Institutional Animal Care and Use Committee (IACUC). A summary of all such exceptions is attached to this annual report. In addition to identifying the IACUC approved exceptions, this summary includes a brief explanation of the exceptions, as well as the species and number of animals affected. The attending veterinarian for this research facility has appropriate authority to ensure the provisions of adequate veterinary care and to oversee the adequacy of other aspects of animal care and 4.) CERTIFICATION BY HEADQUARTERS RESEARCH FACILITY OFFICIAL (Chief Executive Officer (C.E.O.) or Legally Responsible Institutional Official (I.O.J) i certify that the above is true, correct, and complete (7 U.S.C. Section 2143).

SIGNATURE OF C.E.O. OR I.O.

(b)(6), (b)(7)c



APHIS FORM 7023 AUG 2009

P0-2-4-09

Column E Explanation

This form is intended as an aid to completing the Column E explanation. It is not an official form and its use is voluntary. Names, addresses, protocols, veterinary care programs, and the like, are not required as part of an explanation. A Column E explanation must be written so as to be understood by lay persons as well as scientists.

	1	. Registration Number	33-V-0004 / 554			*	
				of animals used in this study.			
	3.			of animals used in the study.			
	4	Explain the procedure pro-	ducing pain and/or dis	stress.			
		Animals are given an oral o	dose of Clostridium D	difficile.			
				NOV 2 7	2009	DE .	
						8 m	
5.	Provide scientific justification why pain and/or distress could not be relieved. State methods or means used to determine that pain and/or distress relief would interfere with test results. (For Federally mandated testing, so litem 6 below) This experiment is designed to investigate the effects of toxin variant strains. Although every effort is made to identify and euthanize animals that become ill following the C. difficile infection, there are some animals that die naturally very suddenly without prior evidence of illness or distress, despite frequent observations (twice daily for 2 weeks, then daily until the end of the experiment). Because not all such animals can be identified by us, and we cannot predict with certainty how many animals will become ill, we have placed all of these animals in USDA category E.						
6.	(C	What, if any, federal regulations 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):					
	A	N/A gency	CED				
		J	CFR	. 11 . 1 . 1 . 1 . 1 . 1 . 1 . 1 . 1 .			

6.