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 it displays a valid OMB control number. The valid OMB control number for this information collection is 0579-0036. The time required to complete this information collection is estimated to average .5 hours 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.

OMB APPROVED 0579-0036		
Interagency Report Control No. 0180-DOA-AN		
Fiscal year:	2021	

## UNITED STATES DEPARTMENT OF AGRICULTURE ANIMAL AND PLANT HEALTH INSPECTION SERVICE

## Annual Report of Research Facility Column E Explanation

(TYPE OR PRINT)

This information is required by law (7 U.S.C. 2143 and 9 C.F.R. §2.36). Failure to report according to the regulations can result in an order to cease and desist.	
2. Research Facility Headquarters address	
675 Hoes Lane West, Room 115 Piscataway, NJ 08854	

3. Number of animals used in the study. 4. Species (common name) of animals used in the study. 134

Hamster

5. Explain the procedure producing pain and distress.

For this study hamsters are used as a model for COVID-19 infection with the ultimate goal of development of therapeutics than can be used in human medicine. Because hamsters are inoculated with live virus, they are expected to have respiratory symptoms.

6. Provide the scientific justification for not providing the appropriate anesthetics, analgesics, or tranquilizing drugs during procedures where the animal experienced accompanying pain or distress greater than momentary or slight.

In order to test potential therapeutic agents, hamsters must first develop clinical signs of COVID-19 infection; administration of analgesics could affect disease progression. While it is possible the treatments may improve the outcome of the disease, it is not known prior to experimentation as these are novel agents and respiratory distress could persist.

7. 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):

Agency	CFR	

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 it displays a valid OMB control number. The valid OMB control number for this information collection is 0579-0036. The time required to complete this information collection is estimated to average .5 hours 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.

OMB APPROVED 0579-0036 Interagency Report Control No. 0180-DOA-AN 2021 Fiscal year:

## UNITED STATES DEPARTMENT OF AGRICULTURE ANIMAL AND PLANT HEALTH INSPECTION SERVICE

## Annual Report of Research Facility

	E Explanation TYPE OR PRINT)	
This information is required by law (7 U.S.C. 2143 and 9 C.F.R. §2.36). Failure to report according to the regulations can result in an order to cease and desist.		
1. REGISTRATION NUMBER	2. Research Facility Headquarters address	
22-R-0157	675 Hoes Lane West, Room 115	
	Piscataway, NJ 08854	
3. Number of animals used in the study. 61	4. Species (common name) of animals used in the study.	
	Rabbit	
5. Explain the procedure producing pain an	d distress.	
For this study rabbits are used as a model to treat Mycobacterium tuberculosis (Mtb), a leading disease among humans responsible for more than a million deaths each year. Because rabbits are inoculated with Mtb, they are expected to have respiratory symptoms.		
	providing the appropriate anesthetics, analgesics, here the animal experienced accompanying pain or	

In order to test potential therapeutic agents, rabbits must first develop clinical signs of Mtb infection; administration of analgesics could affect disease progression. While it is possible the treatments may improve the outcome of the disease, it is not known prior to experimentation as these are novel agents and respiratory symptoms could persist.

7. 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):

Agency	CFR