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 2 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.						
This report is required by law and to be subject to penalties			ions can result in an order to cease	e and desist Interagency Report Control No. 0180-DOA-AN	Fiscal Year 2017	
UNITED STATES DEPARTMENT OF AGRICULTURE ANIMAL AND PLANT HEALTH INSPECTION SERVICE			1. REGISTRATIO 71-G-0001			
				ERS RESEARCH FACILITY (Name, address, and USDA, include ZIP Code)	telephone number as	
ANNUAL REPORT OF RESEARCH FACILITY				HARRY K DUPREE		
(TYPE OR PRINT)			P.O. Box 10	P.O. Box 1050		
				STUTTGART, AR 72160		
3. REPORTING FACILITY (In necessary.)	List all locations where animal			experimentation, or held for these purposes. Attac	h additional sheets, if	
		F	ACILITY LOCATIONS (Sites)			
			-	sary, or use APHIS FORM 7023A.)		
A. Animals Covered By The Animal Welfare Regulations	B. Number of animals being bred, conditioned, or held for use in teaching, testing, experiments, research, or surgery but not yet used for such purposes.	C. Number of animals upon which teaching, research, experiments, or tests were conducted involving no pain, distress, or use of pain-relieving drugs.	D. Number of animals upon which experiments, teaching, research, surgery, or tests were conducted involving accompanying pain or distress to the animals and for which appropriate anesthetic, analgesic, or tranquilizing drugs were used.	E. Number of animals upon which teaching, experiments, research, surgery, or tests we conducted involving accompanying pain or distress to the animals and for which the us appropriate anesthetic, analgesic, or tranquilizing drugs would have adversely affected the procedures, results, or interpretation of the teaching, research, experiments, surgery, or tests. (An explanat of the procedures producing pain or distress these animals and the reasons such drugs were not used must be attached to this report.	e of TOTAL NUMBER OF ANIMALS (Cols. C + D + E) s on	
4. Dogs	0	0	0	0	0	
5. Cats	0	0	0	0	0	
6. Guinea Pigs	0	0	0	0	0	
7. Hamsters	0	0	0	0	0	
8. Rabbits	0	0	0	0	0	
9. Non-human Primates	0	0	0	0	0	
10. Sheep	0	0	0	0	0	
11. Pigs	0	0	0	0	0	
12. Other Farm Animals						
13. Other Animals						
ASSURANCE STATEMENTS	S				· · · · · · · · · · · · · · · · · · ·	

1.) 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.

3.) 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.

4.) 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 use.

CERTIFICATION BY HEADQUARTERS RESEARCH FACILITY OFFICIAL					
(Chief Executive Officer (C.E.O.) or Legally Responsible Institutional Official (I.O.))					
I certify that the above is true, correct, and complete (7 U.S.C. Section 2143).					

DATE SIGNED 13-FEB-2018

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CONTINUATION SHEET FOR ANNUAL REPORT OF RESEARCH FACILITY EXPLANATION FOR APHIS FORM 7023 COLUMN E ENTRIES

FY 2017 CUSTOMER NUMBER: 334026 REGISTRATION NUMBER: 71-G-0001

Species: Channel Catfish Number: 420

Species: Hybrid Striped Bass Number: 1050

Species: Golden Shiner Number: 840

Species: Rainbow Trout Number: 1680

Acute toxicity studies are designed to determine the calculated concentration of a specific test article that will cause 50% mortality in a representative fish population. They are needed to determine the hazardous potential of different compounds and to compare the sensitivity of various fish species that are exposed under distinct environmental conditions. These evaluations must be conducted in a complete living animal in order to understand the complex physiologic and metabolic interactions that contribute to an intoxicated state. A complete assessment of the toxic potential of a test substance cannot currently be accomplished exclusively through available alternative methods. All fish were carefully monitored throughout the study, but some were expected to develop lethargy, loss of appetite, irritated skin or gills, respiratory distress, or aberrant locomotion leading to a moribund condition or death. Humane endpoints were established to minimize pain and discomfort; however, the use of anesthetics, analgesics, or tranquilizing drugs was contraindicated as their administration would have interfered with sensitive metabolic processes and masked clinical and physiological changes that were critical to an accurate assessment of the exposure. The subpopulation of fish that exceeded those parameters is being reported in Column E.

Note: No exceptions to the regulations and standards were requested by the PI or approved by the IACUC.

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CONTINUATION SHEET FOR ANNUAL REPORT OF RESEARCH FACILITY EXPLANATION FOR APHIS FORM 7023 COLUMN E ENTRIES

FY 2017 CUSTOMER NUMBER: 334026 REGISTRATION NUMBER: 71-G-0001

Species: Channel Catfish Number: 794

Species: Hybrid Striped Bass Number: 420

Species: White Bass Number: 420

Fish were used to determine the effectiveness of 1) treatments, 2) immunization (following disease challenge), or 3) diet manipulations following challenge with a bacterial pathogen. Complete living animals were required for these studies to better understand each intervention's impact on an animal's ability to resist and/or respond to infection. The full range of immunologic and physiologic responses to a disease cannot be replicated in any of the *in vitro* systems that are currently available. Some fish may have experienced difficult respiration or changes in the cutaneous osmoregulatory process following bacterial challenge or due to the administration of novel test article. However, anesthetics, analgesics, or tranquilizing drugs could not be given to alleviate these signs as their administration would have interfered with sensitive metabolic processes and masked clinical and physiological changes that were critical to an accurate assessment of the outcome. Humane endpoints were established to minimize pain and discomfort, and fish identified as moribund were euthanized. Some fish may have died before euthanasia could be performed, and this subpopulation is being reported in Column E.

Note: No exceptions to the regulations and standards were requested by the PI or approved by the IACUC.
