

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:	2022
UNITED STATES DEPARTMENT OF AGRICULTURE ANIMAL AND PLANT HEALTH INSPECTION SERVICE Annual Report of Research Facility Column E Explanation <i>(TYPE OR PRINT)</i>			
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	
71-G-001		HARRY K DUPREE P.O. Box 1050 STUTT GART, AR 72160	
3. Number of animals used in the study.		4. Species (common name) of animals used in the study.	
200		ARS Fish	
5. Explain the procedure producing pain and distress.			
Fish will be used to determine the effect of 1) the administration of therapeutics, 2) immunization, 3) genetic strains of dietary feed grains, 4) full/half-sib genetic families, 5) second generation backcross hybrid fish, or 6) antimicrobial feed additives on their health during use of a bacterial disease challenge model. During the bacterial disease model some fish may experience some disruption of respiratory or osmoregulatory processes (skin).			
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.			
Drug intervention (beyond the testing paradigm) is contraindicated because of the scientific need to test the efficacy of therapeutics or immunization strategies in the disease model.			
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):			
None			
Agency			CFR