

<p>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.</p>		<p>OMB APPROVED 0579-0036</p>
		<p>Interagency Report Control No. 0180-DOA-AN</p>
		<p>Fiscal year: 2021</p>
<p>UNITED STATES DEPARTMENT OF AGRICULTURE ANIMAL AND PLANT HEALTH INSPECTION SERVICE</p> <p><b>Annual Report of Research Facility</b> <b>Column E Explanation</b> (TYPE OR PRINT)</p>		
<p>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.</p>		
<p>1. REGISTRATION NUMBER 93-R-0066</p>	<p>2. Research Facility Headquarters address Salk Institute for Biological Studies 10010 N. Torrey Pines Road La Jolla, CA 92037</p>	
<p>3. Number of animals used in the study. 15</p>	<p>4. Species (common name) of animals used in the study. Rabbit</p>	
<p>5. Explain the procedure producing pain and distress.</p> <p>An investigator using rabbits was granted Salk IACUC approval to utilize complete Freund's adjuvant in order to elicit high affinity and avidity antibodies against inoculated epitopes. Animals were given analgesics at the time of antigen/CFA injection, however continuous pain control following injection may not be achieved when analgesic side-effects become a risk (buprenorphine and gastrointestinal stasis; NSAIDS and gastrointestinal ulceration and potential renal impairment), and residual inflammation may lead to further discomfort/pain. These animals are therefore considered category E per the animal use protocol.</p>		
<p>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.</p> <p>Animals were given analgesics at the time of antigen/CFA injection, however continuous pain control following injection may not be achieved when analgesic side-effects become a risk (buprenorphine and gastrointestinal stasis; NSAIDS and gastrointestinal ulceration and potential renal impairment).</p>		
<p>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):</p>		
<p>Agency</p>		<p>CFR</p>