10 F0002

Report to Accompany APHIS Form 7023 to support Column E listing (Unrelieved Pain or Distress) WRAIR, Washington, DC 20307-5100

Pigs -

12/03/99

The lethal shock that is induced by the lethal SE-challenge with the LD50 test in the positive control animals will necessarily cause pain to these animals. But positive controls are required to validate results. Analgesics would impact the physiological parameters, exacerbating the lethal shock or emesis induced by the SE and compromising analysis of collected data. If the experimental drugs proved their utility, the animals should experience relief, but should they not experience relief then that indicates failure of the drug and is necessary for that reason. In all circumstances, the animals will be under constant veterinary care and will not be subject to any unnecessary pain.

Non-human Primates -

In humans, virulent S. flexneri 2a 3457T induces painful abdominal cramps and tenesmus accompanied by myalgia and fatigue. In general, unless the contrary is known or established, it should be assumed that procedures that cause pain in humans also cause pain in animals. Previous studies have shown that opioid down-regulation on the immune response. The potential immunomodulatory effects of buprenorphine have not been evaluated in the NHP model. A study to evaluate buprenorphine will be conducted along with the protocol. If it is not found to alter the specific immunological parameters as measured in this study, then alternatives methods for controlling pain can be deployed. Shigellosis is usually a self-limited infection in rhesus monkeys, and spontaneous recovery is expected after 48 to 72 hours of acute disease. Animals were continually evaluated by the veterinary staff. Guidelines were established to determine progresssion of illness. Once a certain point had been reached, supportive therapy was initiated.

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POC:

Walter Reed Army Institute of Research

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UNITED STATES DEPARTMENT OF AGRICULTURE ANIMAL AND PLANT HEALTH INSPECTION SERVICE 1. REGISTRATION NO. 10-F-003 10 F 000 3 FORM APPROVED OMB NO. 0579-0036

ANNUAL REPORT OF RESEARCH FACILITY

(TYPE OR PRINT)

2. HEADQUARTERS RESEARCH FACILITY (Name and Address, as registered with USDA.

Armed Forces Institute of Pathology Division of Laboratory Animal Medicine 14th & Alaska NW, Bldg 54 Washington DC 20306-6000

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

FACILITY LOCATIONS (Sites)

AFIP Bldg 54, 5th Floor

440

sheets if necessary.)

DEC 0 1 1999

A. Animals Covered By The Animal Wellare 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 were conducted involving accompanying pain or distress to the animals and for which the use of appropriate anesthetic, analgesic, or tranquilizing drugs would have adversely affected the procedures, results, or interpretation of the teaching, research, experiments, surgery, or tests. (An explanation of the procedures producing pain or distress in these animals and the reasons such drugs were not used must be attached to this report).	F. TOTAL NO OF ANIMALS (Cols. C + D + E)
4. Dogs			6		6
5. Cats		_			
6. Guinea Pigs					
7. Hamsters					
8. Rabbits	5	4	280		28 9
9. Non-human Primates					
10. Sheep		·			
11. Pigs			6		6
12. Other Farm Animals					
13. Other Animals					
Mice		2789	13		2802
Rat	50	150	88		288
					•

- 1) Prolessionally acceptable standards governing the care, treatment, and use of animals, including approriate 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 provision of adequate veterinary care and to oversee the adequacy of other aspects of

CERTIFICATION BY HEADQUARTES RESEARCH FACILITY OFFICIAL (Chief Executive Officer or Legally Responsible Institutional Official)

Licertify that the above is true, correct, and complete (7 U.S.C. Section 2143)

NAME & TITLE OF C.E.O. OR INSTITUTIONAL OFFICIAL (Type or Print)

DATE SIGNED

(AGG 91)

(Replace VS FORM 18-23 (OCT 88), which is obsolete)

USN