FACILITY LOCATIONS (Sites) - See Atached Listing

A. Animals Covered By The Animal Welfare Regulations	B. Number of animal being bred, conditioned, or held for use in teaching, testing, experiments, research, or surgery but not ye used for such purposes.	C. Number of animals upon which teaching, research, experiments, or tests were conducted involving no pain, distress, or use or 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 an for which appropriate anesthetic, analgesic, or tranquilizing drugs were used.	E. Number of animats 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 tranquilized rugs would have adversely affected the procedures, resear interpretation of the teaching, research, experiments, surgery, or tests. (An explanation of the procedures producing pain or distress in these animals and the reason such drugs were not used must be attached to this report	F. TOTAL NUMBE: OF ANIMALS (COLUMNS C + D + E)
4. Dogs					51F
5. Cats		. <u>.</u>	-		**
6. Guinea Pigs	-	1499		326	1825
7. Hamsters	-	<u> </u>		320	1023
8. Rabbits		1500			1500
9. Non-human Primates					1300
t0. Sheep			-		
11. Pigs		<u>-</u>			
12. Other Farm Animals	,				****
3. Other Animals					1411
**					

SSURANCE STATEMENTS

- 1) Professionally acceptable standards governing the care, treatment, and use of animals, including appropriate use of anestetic, analgesic, and tranquilizing drugs, prior to, during, and following actual researching, testing, surgery, or experimentation were followed by this research facility.
- 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 ap 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 included in 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 animal care and use.

	CERTIFICATION BY HEADQUARTERS RESEARCH FACILITY OFFICIAL (Chief Executive Officer or Legally Responsible Institutional Official)									
SIGNATURE	ECEO OR INSTITUTIONAL OFF	L.	NAME & TITLE OF C.E.O. OR INSTITUTIONAL OFFICIAL (Type or Print)	DATE SIGNED						
APHIS FOR	(b)(6), (b)(7)c	(OCT 88), which is obsolete.)	(b)(6), (b)(7)c	11/05/08						

Column E Explanation

This form is intended as an aid to completing the Column E explanation. It is not an official form and its use is voluntary. Names, addresses, protocols, veterinary care programs, and the like, are not required as part of an explanation. A Column E explanation must be written so as to be understood by lay persons as well as scientists.

	1.	Registration Number: 5	R-0127					
2	2.	2. Number326	of animals used in this study.					
3	3,	3. Species (common name) <u>Gui</u>	ea_pigsof animals used in the study.					
4	4. Explain the procedure producing pain and/or distress.							
		Tetanus potency - To o	anitate the potency of Tetanus Immune Globulin (Huma	n)				
		by using neutralization	methodology.					
5		Provide scientific justification whe determine that pain and/or distress tem 6 below)	pain and/or distress could not be relieved. State methods or means used to s relief would interfere with test results. (For Federally mandated testing, s	o ee				
		This testing is manda Administration of ane with the outcome of t	ed by Talecris Product License Testing requirements. Thetic, analgesic or tranquilizer drugs could interfe t testing.	ere!				
6.	V (1	What, if any, federal regulations (CFR) title number and the speci	quire this procedure? Cite the agency, the code of Federal Regulations section number (e.g., APHIS, 9 CFR 113.102):					
	Α	Agency FDA	CFR 21CFR 610.10 USP 29(2006) page 2090					