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 0579-0013. The time required to complete this information collection is estimated to average 1 hour 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 application must be submitted for issuance of a United States Veterinary Biological Product Permit. This information will be used to determine if the product may be brought into the United States, or for approval of transit shipment of biological products move through the United States (9 CFR 104). INSTRUCTIONS: Submit an application for each product. If more space is needed, attach additional sheets and refer to block number. Enclose supporting documents.

UNITED STATES DEPARTMENT OF AGRICULTURE

USDA PERMITTEE NUMBER (LEAVE BLANK FOR INITIAL APPLICATIONS).

ANIMAL AND PLANT HEALTH INSPECTION SERVICE VETERINARY SERVICES, CENTER FOR VETERINARY BIOLOGIC APPLICATION FOR			DATE SUBMITTED			
						UNITED STATES VETERINARY BIOL
2. TYPE OF APPLICATION						
RESEARCH AND EVALUATION (Complete all items except 10 through 15)						
3. NAME AND ADDRESS OF APPLICANT (Include Number, Street	t or RFD Number, City, State, and ZIP	Code) 4. NAME	AND ADDRESS OF PRO	DUCER		
5.NAME OF PRODUCT (one only)	T	EOD EVOR 8	HIPMENT OF SAME PRO	DUCT GIVE		
S.NAME OF PROBOCT (UND UNITY)	6. ESTIMATED ARRIVAL DATE		ATED QUANTITY		STATES PORT OF ENTRY	
9.IF PRODUCT IS FOR RESEARCH AND EVALUATION, FURNIS methods of propagation including composition of medium; species	H NAME AND LOCATION OF INSTITU	JTION DOING RE	SEARCH (If same as Item	n 3, so state. Enclos	se brief description of product;	
pursuant to 9 CFR 104.4(a).)	or animais or cen cultures used, method	u oi illactivation o	alleridation, recommende	allori for use, and pr	oposeu piari oi evaluation	
10.IF PRODUCT FOR GENERAL DISTRIBUTION AND SALE (End	close manufacturer's or producer's agre	eement regarding	preparation testing and l	abeling of products	and inspection facilities	
Enclose supporting documents specified in 9 CFR 104.5.))			p - p			
11.ADDRESS OF STORAGE FACILITIES (If different from Item 3)			2. TYPE OF ORGANIZATION			
•			CORPORATION PARTNERSHIP INDIVIDUAL			
		40.15.00	<u>-</u>			
			rticles of Incorporation)	E IN WHICH INCO	RPORATED (Enclosed certified	
	14. PRINCIPAL OFFICER	S OR PARTNERS	3			
A. NAME OF EACH	B. TITLE		C. BUSINESS ADDRESS (Include Number and Street, or RFD Number, City, State, and ZIP Code)			
			(Include Number a	and Street, or RFD I	vurriber, City, State, and ZIF Code)	
A. DESTINATION		15.IF TRANSIT SHIPMENT GIVE B. CARRIER(S)		C SCHEDIII E	(Dates in transit)	
A. DESTINATION	b. CARRIER(3)		Arrival	C. SCHEDULE	Departure Departure	
	CERTIFICA		I		1	
In accordance with the Act of Congress approved March 4 biological product for the purpose specified in item 2 abov						
regulations and orders of the Department governing the in						
deceive in any particular. 16. SIGNATURE OF AUTHORIZED OFFICIAL		17. TITLE			B. DATE SIGNED	