UNITED STATES DEPARTMENT OF AGRICULTURE ANIMAL AND PLANT HEALTH INSPECTION SERVICE VETERINARY BROLOGICS 1920 DAYTON AVENUE AMES, IOWA 50010     ADVERSE EVENT REPORT     ADVERSE EVENT REPORT     Image: Services Service Service Service Service Service Service Service Advectment Service
1920 DAYTON AVENUE AMES, IOWA 50010     See reverse side for Privacy Act notice     1. Information Reported By
1. Information Reported By   Clinical Pathology Laboratory   Distributer   Human Patient     Medical Physician   Owner/Producer/Employee   Other     2. First Name   3. Last Name   4. Contact Number   5. Submitter's Case Number     6. Date First Received (MM/DD/YYYY)   7. Report   8. Submitted to Manufacturer   9. Country of Occurrence     10. Case Type   Animal Complaint   Human Exposure   Product Problem Only     11. Problem Type   Ecc-toxicity   Extra Label Use   Human Exposure - Asymptomatic     PRODUCT INFORMATION   Product Problem   Generic Name/Active Ingredient(s)
Attending Veterinarian   Clinical Pathology Laboratory   Distributer   Human Patient     Medical Physician   Owner/Producer/Employee   Other   Submitted's Case Number     2. First Name   3. Last Name   4. Contact Number   5. Submitter's Case Number     6. Date First Received   7. Report   8. Submitted to Manufacturer   9. Country of Occurrence     (MMDD/YYY)   Initial   Follow-up   Yes   No     10. Case Type   Animal Complaint   Human Exposure   Product Problem Only     11. Problem Type   Ecc-toxicity   Extra Label Use   Human Exposure - Asymptomatic     Human Exposure - Symptomatic   Lack of Efficacy   Product Problem     PRODUCT INFORMATION
Medical Physician   Owner/Producer/Employee   Other     2. First Name   3. Last Name   4. Contact Number   5. Submitter's Case Number     6. Date First Received   7. Report   8. Submitted to Manufacturer   9. Country of Occurrence     (MM/DD/YYYY)   Initial   Follow-up   Yes   No     10. Case Type   Initial   Follow-up   Product Problem Only     11. Problem Type   Adverse Reaction   Eco-toxicity   Extra Label Use   Human Exposure - Asymptomatic     Human Exposure -   Lack of Efficacy   Product Problem   Product Problem     PRODUCT INFORMATION
2. First Name   3. Last Name   4. Contact Number   5. Submitter's Case Number     6. Date First Received (MM/DD/YYYY)   7. Report   8. Submitted to Manufacturer   9. Country of Occurrence     10. Case Type   Initial   Follow-up   Yes   No   9. Country of Occurrence     10. Case Type   Animal Complaint   Human Exposure   Product Problem Only   9. Country of Occurrence     11. Problem Type   Adverse Reaction   Ecc-toxicity   Extra Label Use   Human Exposure - Asymptomatic     Human Exposure -   Lack of Efficacy   Product Problem   Product Problem     PRODUCT INFORMATION     Product   Brand Name/Trade Name   Generic Name/Active Ingredient(s)
6. Date First Received (MM/DD/YYYY)   7. Report   8. Submitted to Manufacturer   9. Country of Occurrence     10. Case Type   Initial   Follow-up   Yes   No     10. Case Type   Animal Complaint   Human Exposure   Product Problem Only     11. Problem Type   Adverse Reaction   Eco-toxicity   Extra Label Use   Human Exposure - Asymptomatic     Human Exposure - Symptomatic   Lack of Efficacy   Product Problem     PRODUCT INFORMATION     Product Number   Brand Name/Trade Name
(MM/DD/YYYY)   Initial   Follow-up   Yes   No     10. Case Type   Initial   Follow-up   Yes   No     10. Case Type   Initial   Human Exposure   Product Problem Only     11. Problem Type   Initial   Ecc-toxicity   Extra Label Use   Human Exposure - Asymptomatic     Image: Human Exposure - Symptomatic   Initial   Ecc-toxicity   Product Problem   Human Exposure - Asymptomatic     PRODUCT INFORMATION     Product Name/Trade Name     Generic Name/Active Ingredient(s)
Initial   Follow-up   Yes   No     10. Case Type   Animal Complaint   Human Exposure   Product Problem Only     11. Problem Type   Adverse Reaction   Ecco-toxicity   Extra Label Use   Human Exposure - Asymptomatic     Human Exposure - Symptomatic   Lack of Efficacy   Product Problem   Asymptomatic     Product   Brand Name/Trade Name   Generic Name/Active Ingredient(s)
Animal Complaint   Human Exposure   Product Problem Only     11. Problem Type   Adverse Reaction   Ecc-toxicity   Extra Label Use   Human Exposure - Asymptomatic     Human Exposure - Symptomatic   Lack of Efficacy   Product Problem   Asymptomatic     PRODUCT INFORMATION     Product   Brand Name/Trade Name   Generic Name/Active Ingredient(s)
11. Problem Type     Adverse Reaction   Eco-toxicity     Human Exposure -   Asymptomatic     Human Exposure -   Lack of Efficacy     Product Problem   Product Problem     Product Number   Brand Name/Trade Name     Generic Name/Active Ingredient(s)
Adverse Reaction   Eco-toxicity   Extra Label Use   Human Exposure - Asymptomatic     Human Exposure - Symptomatic   Lack of Efficacy   Product Problem     PRODUCT INFORMATION     Product Number     Brand Name/Trade Name   Generic Name/Active Ingredient(s)
Adverse Reaction Eco-toxicity Extra Label Ose Asymptomatic   Human Exposure - Symptomatic Lack of Efficacy Product Problem   PRODUCT INFORMATION   Product Number Brand Name/Trade Name Generic Name/Active Ingredient(s)
Symptomatic Lack of Efficacy Product Problem   PRODUCT INFORMATION   Product Number Brand Name/Trade Name Generic Name/Active Ingredient(s)
Product     Brand Name/Trade Name     Generic Name/Active Ingredient(s)
Number Brand Name/Trade Name Generic Name/Active Ingredient(s)
1
2
3
4
Product 1     Product 2     Product 3     Product 4
Manufacturer
Serial/Lot Number
Expiration Date
Was product used as per Yes No Not Applicable Yes No Not Applicable Yes No Not Applicable Yes No Not Applicable
Iabel instructions?     Unknown to Reporter     Unknown to Reporter     Unknown to Reporter     Unknown to Reporter
Off-label use type
Has patient received this Yes No Not Applicable Yes No Not Applicable Yes No Not Applicable Yes No Not Applicable
product before Unknown to Reporter Unknown to Reporter Unknown to Reporter Unknown to Reporter
Has patient experienced   Yes   No   Not Applicable   Yes   No   Not Applicable   Yes   No   Not Applicable     AEs from this product   Yes   No   Not Applicable   Yes   No   Not Applicable   Yes   No   Not Applicable
before? Unknown to Reporter Unknown to Reporter Unknown to Reporter Unknown to Reporter
Route of Administration
Site of Administration
Site of Administration Start Date End Date Start Date End Date End Date Start Date End Date End Date   Duration of Treatment/Exposure Start Date End Date Start Date End Date Start Date End Date End Date
Duration of Start Date End Date Start Date End Date Start Date End Date
Duration of Treatment/Exposure Start Date End Date Start Date End Date Start Date End Date

DETAILED DESCRIPTION OF EVENT (narrative)									
Event Category									
Anaphylaxis - Hypersensitivity	Autoimmune	Birth Defect	Lack of Expected Efficacy						
Local	Neoplasia	Reproductive	Other						
What was the final outcome?									
Alive with Sequelae	Death (All Causes)	🗌 Euthanasia	Natural Death						
Recovered	Remains Under Treatment	Not Applicable	Unknown						
Enter case narrative (if necessary continue on page 3):									

Enter case narrative (if necessary, continue on page 3):

SUSPECTED ADVERSE EVENT DATE(S)									
1. Date of Onset of	1. Date of Onset of AE (MM/DD/YYYY)		2. Duration of Susp	ation of Suspected Adverse Event 3. Time Between Adn		n Administration an	d Event		
1. Number of Animals Exposed		2. Number of Anima		3. Number of D	3. Number of Dead Animals				
4. Animal Condition Prior to Treatment									
5. Animal Name		Fair Good			Poor Not Applicable Unknow		Jown		
		Fema		e 🗌 Mixed	Not Applicable 🔲 Unknown		nknown		
7. Species	Cattle	Chicke	en 🗌 Dog	Goat	Horse	] Human	Other		
	Mixed with		9. Status				outor		
			Intact	Neutered	🗌 Not App	🗌 Not Applicable 🗌 Unknown			
10. Age From		11. Age To		12. Weight From	13. \	Weight To			
	I.		REPORTER	INFORMATION					
Primary Report									
1. Sender	Oliniaal D					an/Draducent			
Attending Veterinaria	an Clinical Pa		Distributor	Human Patient III		ner/Producer/	Other		
2. First Name 3. Last Name									
4. Address (include ZIP Code and country)									
5. Phone Number 6. Fax Nun		nber	er 7.Email						
Additional Informat	ion								
Additional mormat	1011								
Save and submit vi	ia email to:	F	Print form and mail to:		Print and fax it t	Print and fax it to:			
			Pharmacovigilance, USDA,						
cvb@aphis.usda.com		Center for Veterinary Biologics, 1920 Dayton Avenue,		515-337-6120					
		Ames, IA 50010							
PRIVACY ACT NOTICE The information requested on this form will not be retrieved from our files by using your name or personal identifier and is therefore, in the									
opinion of this a				1974. However, in keepi					
Authority:	9 CFR Section 11	4.7.							
Purpose:	That compliance with the Act and applicable regulations be under supervision of person(s) competent in the preparation of biological products.								

To determine that the responsible person(s) producing biological products are qualified by training and experience and have demonstrated fitness to produce such products in compliance with the Act. Routine uses:

## **CONTINUATION SHEET**

(use this page to continue any item on this form)