DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION

DISTRICT ADDRESS AND PHONE NUMBER	DATE(S) OF INSPECTION
404 BNA Dr., Bldg. 200, Ste. 500 Nashville, TN 37217-2597 (615)366-7801 Fax:(615)366-7802	4/27/2026-5/1/2026
	FEI NUMBER 3038121665

NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED
Matthew R. Jacobs, Vice President

FIRM NAME	STREET ADDRESS
RAMVet LLC	100 Ponchatoula Pkwy
CITY, STATE, ZIP CODE, COUNTRY	TYPE ESTABLISHMENT INSPECTED
Ponchatoula, LA 70454-8309	Animal Drug Manufacturer

This document lists observations made by the FDA representative(s) during the inspection of your facility. They are inspectional observations, and do not represent a final Agency determination regarding your compliance. If you have an objection regarding an observation, or have implemented, or plan to implement, corrective action in response to an observation, you may discuss the objection or action with the FDA representative(s) during the inspection or submit this information to FDA at the address above. If you have any questions, please contact FDA at the phone number and address above.

DURING AN INSPECTION OF YOUR FIRM I OBSERVED:

OBSERVATION 1

Written procedures are not established for the cleaning and maintenance of equipment, including utensils, used in the manufacture, processing, packing or holding of a drug product.

Specifically, Your firm failed to establish a validated cleaning procedure for equipment used in the manufacture and filling of (b) (4) (b) (4) Solution) (b) (4) . Your firm has not demonstrated that the cleaning procedures for the (b) (4) Gallon Mix Tank (b) (4) Gallon Mix Tank Agitator ((b) (4), and (b) (4) Weigh Tank ((b) (4)) adequately remove residues of (b) (4) , and/or cleaning agents to predetermined, scientifically justified acceptance limits.

SEE REVERSE OF THIS PAGE

EMPLOYEE(S) SIGNATURE
Demario L Walls, Investigator

X Demario L Walls
Investigator
Signed By: DEMARIO L. WALLS - S
Date Signed: 05-01-2026 10:08:18

DATE ISSUED
5/1/2026

The observations of objectionable conditions and practices listed on the front of this form are reported:

1. Pursuant to Section 704(b) of the Federal Food, Drug and Cosmetic Act, or
2. To assist firms inspected in complying with the Acts and regulations enforced by the Food and Drug Administration.

Section 704(b) of the Federal Food, Drug, and Cosmetic Act (21 USC 374(b)) provides:

"Upon completion of any such inspection of a factory, warehouse, consulting laboratory, or other establishment, and prior to leaving the premises, the officer or employee making the inspection shall give to the owner, operator, or agent in charge a report in writing setting forth any conditions or practices observed by him which, in his judgment, indicate that any food, drug, device, or cosmetic in such establishment (1) consists in whole or in part of any filthy, putrid, or decomposed substance, or (2) has been prepared, packed, or held under insanitary conditions whereby it may have become contaminated with filth, or whereby it may have been rendered injurious to health. A copy of such report shall be sent promptly to the Secretary."