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	3/31/2026-4/7/2026* FEI NUMBER 2313447

NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED
B. Karen Payne, Quality Program Supervisor

FIRM NAME	STREET ADDRESS
Purina Animal Nutrition LLC	64080 Highway 51
CITY, STATE, ZIP CODE, COUNTRY	TYPE ESTABLISHMENT INSPECTED
Roseland, LA 70456-1926	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

You did not identify and implement preventive controls to ensure that any hazards requiring a preventive control are significantly minimized or prevented.

Specifically, your Food Safety Plan does not identify aflatoxin, fumonisin, and vomitoxin (DON) as a hazard that requires a preventative control. On (b) (4) you manufactured (b) (4) bags of Purina Nature's Match Sow & Pig Complete (Lo (b) (4) , which were subsequently (b) (4)
(b) (4)

*DATES OF INSPECTION

3/31/2026(Tue), 4/01/2026(Wed), 4/07/2026(Tue)

SEE REVERSE OF THIS PAGE	EMPLOYEE(S) SIGNATURE		DATE ISSUED
	Sidney M Smith, Investigator	X Sidney M Smith Investigator Signed By: SIDNEY M. SMITH -S Date Signed: 04-07-2026 08:55:49	4/7/2026

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."