DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION

DISTRICT ADDRESS AND PHONE NUMBER	DATE(S) OF INSPECTION
8050 Marshall Drive, Suite 205 Lenexa, KS 66214 (913)495-5100 Fax:(913)495-5115	4/30/2024-5/2/2024
	FEI NUMBER 1924965

NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED
Justin P. Wilkinson, Plant Manager

FIRM NAME	STREET ADDRESS
Nestle Purina Pet Care Co.	2200 Manufacturing Dr
CITY, STATE, ZIP CODE, COUNTRY	TYPE ESTABLISHMENT INSPECTED
Clinton, IA 52732-6846	Pet Food 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 WE OBSERVED:

OBSERVATION 1

You did not conduct a reanalysis of your food safety plan as appropriate.

Specifically, Nestle Purina Pet Care Co. was notified by a representative of the Food and Drug Administration regarding a new potential hazard in pet food manufactured at your facility. As of 04/30/2024, your local management team indicated that they were unaware of this new potential hazard and had not conducted a reanalysis of your food safety plan.

X Joseph R Haynes
Investigator
Signed By: Joseph R. Haynes -S
Date Signed: 05-02-2024 14:32:43

SEE REVERSE OF THIS PAGE

EMPLOYEE(S) SIGNATURE
Andrew K Schaal, Investigator
Joseph R Haynes, Investigator

X Andrew K Schaal
Investigator
Signed By: Andrew K. Schaal -S
Date Signed: 05-02-2024 14:31:23

DATE ISSUED
5/2/2024

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."