23-036-9019

Feb 05, 2023
RECEIVED DATE
Feb 03, 2023

SEND TO **35432**



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REPORT OF ANALYSIS

For: (35432) OPEN FARM - Dog-OF Rawmix Beef

	Level Found		Reporting		Analyst-	Verified-
Analysis	As Receive	d Units	Limit	Method	Date	Date
Sample ID: 033231317 - Micro	Lab Number: 13986056	Date Sampled: 20	023-02-02			
Listeria	negative	e org/125g	1	RapidChek/AOAC RI 020401	kkb0-2023/02/05	jzh4-2023/02/05
E. coli (generic)	n.d	I. cfu/g	10	AOAC OMA 991.14	Jcp2-2023/02/05	jzh4-2023/02/05
Salmonella	negative	e org/375g	1	RapidChek/AOAC RI 030301; AFNOR SDI 34/01-04/10	kkb0-2023/02/05	jzh4-2023/02/05

All results are reported on an AS RECEIVED basis., n.d. = not detected, cfu = colony forming unit

For questions please contact:

Jamie Wood Account Manager

jwood@midwestlabs.com (402)590-2964

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REPORT OF ANALYSIS

Detailed Method Description(s)

E. coli and Total Coliform using 3M Petrifilm

Sample analysis follows MWL MI 292 which is based on AOAC OMA 991.14. A representative sample is obtained and added to phosphate buffer. Aliquots of the sample are withdrawn and placed on Petrifilm plates. The plates are incubated for 44 to 52 hours. After incubation, the plates are counted to determine the number of generic E. coli and total coliforms present. The color of the colony differentiates a generic coliform from E. coli. The levels are reported as colony forming units (cfu).

Salmonella - Lateral Flow

Samples are analyzed following MWL MI 195 which is based on the RapidChek Select Salmonella User Guide. A representative sample is obtained and combined with a selective media and allowed to incubate. After incubation, a test strip is used for Salmonella determination. Results are reported as negative or presumptive positive.

Listeria Lateral Flow

Samples are analyzed following MWL MI 194 which is based on the RapidChek Listeria User Guide. A representative sample is obtained and combined with a selective growth media. It is incubated for 40-48 hours. After incubation, an aliquot is heated for 10 minutes, and a test strip for Listeria detection is used. Results are reported as negative or presumptive positive. This procedure does not speciate Listeria.