

REPORT NUMBER

22-346-9037

REPORT DATE
Dec 12, 2022
RECEIVED DATE
Dec 10, 2022

SEND TO
35432



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ISSUE DATE
Dec 12, 2022

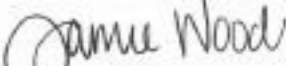
REPORT OF ANALYSIS

For: (35432) OPEN FARM -
341221109

Analysis	Level Found	Reporting			Analyst-Date	Verified-Date
	As Received	Units	Limit	Method		
Sample ID: 341221109 - Micro Lab Number: 13952136 Date Sampled: 2022-12-07						
Listeria	negative	org/125g	1	RapidChek/AOAC RI 020401	Lrs8-2022/12/12	mml4-2022/12/12
E. coli (generic)	n.d.	cfu/g	10	AOAC 2018.13	Jcp2-2022/12/11	snl7-2022/12/11
Salmonella	negative	org/375g	1	RapidChek/AOAC RI 030301; AFNOR SDI 34/01-04/10	Hct6-2022/12/12	mml4-2022/12/12

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

For questions please contact:


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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 2018.13. 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 Aliquots of the sample are withdrawn and placed on Petrifilm plates.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 18 to 24 hours. After incubation, the plates are counted to determine the number of generic E. coli and total coliforms present. The color of the colony and the presence of gas differentiate 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.

The result(s) issued on this report only reflect the analysis of the sample(s) submitted.

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