

**DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION**

DISTRICT OFFICE ADDRESS AND PHONE NUMBER 158-15 Liberty Avenue Jamaica, NY 11433 (718) 340-7000 Ext:5301 Fax: (718) 662-5661	DATE(S) OF INSPECTION 1/30/2026-3/4/2026*
	FEI NUMBER 3000206705
Industry Information: www.fda.gov/oc/industry	

NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED Chad Keeney, Operations Manager	
FIRM NAME Lifetime Brands, Inc.	STREET ADDRESS 1000 Stewart Ave
CITY, STATE, ZIP CODE Garden City, NY 11530-4814	E-MAIL ADDRESS (b) (6), (b) (7)(C)

This document lists observations made by the FDA representative(s) during the review of your Foreign Supplier Verification Program (FSVP). They are 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) including by submitting 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 FOREIGN SUPPLIER VERIFICATION PROGRAM I OBSERVED:

OBSERVATION 1

Your hazard analysis did not identify a hazard that requires a control.

Specifically,

This is a repeat observation from the previous inspection(s) conducted on 12/09/2022 (written observation), 02/04/2022 (written observation).

For Ground Cumin that you imported from (b) (4), Black Peppercorn that you imported from (b) (4) and (b) (4) of Black Peppercorn in each (b) (4) that you imported from (b) (4), your hazard analysis did not identify a hazard that requires a control.

OBSERVATION 2

You did not document that you or another entity reevaluated a foreign supplier's performance and the risk posed by a food during or at the end of a 3-year period.

Specifically,

For Ground Cumin that you imported from (b) (4), you did not document that you or another entity reevaluated a foreign supplier's performance and the risk posed by a food during or at the end of a 3-year period.

SEE REVERSE OF THIS PAGE	EMPLOYEE(S) SIGNATURE Ayorinde O Babalola, Investigator	Ayorinde O Babalola Investigator Signed By: AYORINDE O. BABALOLA -S Date Signed: 03-04-2026 11:36:16 X	DATE ISSUED 3/4/2026

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OBSERVATION 3

You did not establish written procedures for ensuring that appropriate foreign supplier verification activities are conducted with respect to a food you import.

Specifically,

This is a repeat observation from the previous inspection(s) conducted on 12/09/2022 (written observation), 02/04/2022 (written observation).

For Ground Cumin that you imported from (b) (4), Black Peppercorn that you imported from (b) (4) and (b) (4) of Black Peppercorn in each (b) (4) that you imported from (b) (4), you did not establish adequate written procedures for ensuring that appropriate supplier verification activities are conducted with respect to a food you import.

OBSERVATION 4

You did not determine and document the frequency of conducting the verification activities to provide adequate assurances that a food you obtain from a foreign supplier is produced in compliance with processes and procedures that provide the required level of public health protection.

Specifically,

This is a repeat observation from the previous inspection(s) conducted on 12/09/2022 (written observation).

For Ground Cumin that you imported from (b) (4), Black Peppercorn that you imported from (b) (4), and (b) (4) of Black Peppercorn in each

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(b) (4) that you imported from (b) (4), you did not determine and document the frequency of conducting the verification activities to provide adequate assurances that a food you obtain from a foreign supplier is produced in compliance with processes and procedures that provide the required level of public health protection.

OBSERVATION 5

You did not document your review and assessment of a hazard analysis conducted by another entity and that a hazard analysis conducted by another entity was conducted by a qualified individual.

Specifically,

For Ground Cumin that you imported from (b) (4), Black Peppercorn that you imported from (b) (4), you did not document your review and assessment of a hazard analysis conducted by another entity and that a hazard analysis conducted by another entity was conducted by a qualified individual.

***DATES OF INSPECTION**

1/30/2026(Fri), 2/02/2026(Mon), 2/04/2026(Wed), 2/10/2026(Tue), 2/13/2026(Fri), 2/19/2026(Thu), 2/25/2026(Wed), 3/03/2026(Tue), 3/04/2026(Wed)

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	X		

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

1. Pursuant to Section 805 of the Federal Food, Drug and Cosmetic Act, or
2. To assist firms inspected in complying with the laws and regulations enforced by the Food and Drug Administration.