

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

DISTRICT ADDRESS AND PHONE NUMBER 404 BNA Dr., Bldg. 200, Ste. 500 Nashville, TN 37217-2597 (615) 366-7801 Fax: (615) 366-7802 Industry Information: www.fda.gov/oc/industry		DATE(S) OF INSPECTION 05/31/2011 - 06/02/2011
NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED TO: Robert Wilford Webb, III, Director of Production		FEI NUMBER 3004327090

FIRM NAME Cal-Maine Farms, Inc.	STREET ADDRESS 3280 Adams Lane
CITY, STATE, ZIP CODE, COUNTRY Edwards, MS 39066	TYPE ESTABLISHMENT INSPECTED 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

Your written SE prevention plan lacks appropriate SE prevention measures.

Specifically, your firm failed to have written procedures that address egg testing for Salmonella Enteritidis following a positive environmental test of the pullet environment.

SEE REVERSE OF THIS PAGE	EMPLOYEE(S) SIGNATURE Amanda K. Willey, Investigator <i>Amanda K. Willey</i> Sidney M. Smith, Investigator <i>Sidney M. Smith</i> Wayne S. Fortenberry, Investigator <i>Wayne S. Fortenberry</i>	DATE ISSUED 06/02/2011
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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 judgement, 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."