

Gary W. Black, Commissioner
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Recall Guidance for Industry in Georgia

The impact of a recall on your business can be minimized by having an efficient, organized and expedited response. In cooperation with the Georgia Department of Agriculture (GDA), U.S. Food and Drug Administration (FDA) and/or U.S. Department of Agriculture Food Safety Inspection Service (USDA-FSIS), you will be able to work through the recall process. This document is intended to provide general guidance on executing and recovering from a recall. Several steps may take place concurrently.

Important Contacts:

- Natalie Adan, GDA Food Safety Director, 404-657-4801, Natalie.Adan@agr.georgia.gov
- Craig Nielsen, GDA Food Safety Deputy Director, 404-656-3627, Craig.Neilsen@agr.georgia.gov
- Lt. James Betz, MPH, CPH, State Liaison, Atlanta District, 843-746-2990 x19, james.betz@fda.hhs.gov
- Emma Nesbit, Recall Coordinator, Atlanta District, 404-253-1293, orahafeast3recalls@fda.hhs.gov
- Geneva Dennis, FSIS District Case Specialist, Atlanta District, 404-562-5900, Geneva.Dennis@fsis.usda.gov

Reporting the Recall:

□ Within the first 24 hours: If a finished product/ingredient sample has tested positive (even if the product is still in your control) contact the GDA at 404-657-4801or send details to positiveresults@agr.georgia.gov. ○ Complete a Reportable Food Registry (RFR) using your facility's food registry number online at www.fda.gov/food/complianceenforcement/rfr/. Some fields may be left blank initially and amended later as more information becomes available. For help, contact RFRSupport@fda.hhs.gov. □ Make direct contact with the GDA, FDA and, if needed, the USDA-FSIS Atlanta District. Be prepared to provide information about the status of your product, nature of the recall, root cause behind the issue (if known), any consumer complaints, distribution information, etc. □ The Recall Coordinator with FDA and/or GDA will support your recall operations and provide assistance in drafting a public notification about the recall. Things to include: Product descriptions, distribution, sizes, lot codes, types of packaging, etc. □ The GDA and FDA will determine primary jurisdiction at your facility and will plan to conduct an onsite recall investigation.

The Recall Investigation:

Prompt identification of affected product(s) and source(s) of contamination means a reduced scope of the recall and time to product withdraw. Helping your regulatory officials in turn helps ensure an efficient and effective recall.

- Help the federal and state inspectors obtain the following:
 - Scope of the recall, which ensures the recall is not too broad (leading to additional losses) or too narrow (requiring subsequent or expanded recalls).
 - O Up-to-date inventory for finished product, in-process material and/or raw materials present in your facility.

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0	Up-to-date distribution, including a list of customers who received the product (could be sales
	records, shipping manifests, invoices or other business documents). This information is classified as
	proprietary and is protected from disclosure.

Provide assistance and education to GDA & FDA about the meaning of product or lot codes used in
your facility. The more rapidly specific products or lots affected are catalogued, the more rapidly the
recall can be concluded.

Public Notification:

Your public notification campaign will be an integral part of the product correction/withdraw. Prompt,
forthcoming communication with the public will protect your business and hasten the recovery process.
☐ Work with regulatory Recall Coordinator(s) to draft a recall press release notifying the public.
☐ Include as much information as possible about:
 Product name
o Packaging description size LIPC and/or lot code information

- Packaging description, size, UPC and/or lot code information
- o The reason behind the recall, including the contaminant of concern
- Areas of distribution
- o Disposal instructions and availability for refunds
- □ While GDA, FDA and/or USDA may issue their own press release, every opportunity will be given to allow the firm to release the information first.

Restoration and Recovery:

Once the recall is initiated, begin planning your restoration, recovery and strategy to prevent future occurrences. The GDA will provide support, approval and verification for your restoration and recovery process. FDA and USDA will also be available during this phase and throughout the recall process.