RECALL NOTICE

The following message is being sent to all authorized member recall personnel.

Updated information is now available. The lists of retail consignees have been posted for recalls:

- 069-2019, Ruiz Foods Products Inc. Recalls Bacon Breakfast Wrap Products Due to Possible Foreign Matter Contamination (Jun 14, 2019)
- 070-2019, C&S Wholesale Grocers Recalls Meat and Poultry Products due to Possible Temperature Abuse During Transport at Two Stores in New York (Jun 20, 2019)

*** We believe that the products being recalled were not processed or offered through the national office.**
* We strongly encourage you to notify your agencies within one business day from receipt of this notice.

* ALL cased and uncased inventories, both at the member level and agency level, need to be checked. This product may have entered member and agency warehouses through salvage, local donations, TEFAP, local purchases, retail pickups, food drives, or other avenues.

* For additional local details, please contact the Health Department(s) for the area(s) your food bank serves.

RECALL NOTICE

The following message is being sent to all authorized member recall personnel.

News Release

San Giuseppe Salami Co. by Giacomo Recalls Ready-To-Eat, Frozen Andouille Sausage Products due to Possible Foreign Matter Contamination

Class I Recall 071-2019
Health Risk: High Jun 24, 2019

Congressional and Public Affairs
Felicia Thompson

(202) 720-9113
Press@fsis.usda.gov

WASHINGTON, June 24, 2019 – San Giuseppe Salami Co. by Giacomo, an Elon, N.C., establishment, is recalling approximately 832 pounds of ready-to-eat (RTE), frozen andouille sausage products that may be contaminated with extraneous materials, specifically metal, the U.S. Department of Agriculture's Food Safety and Inspection Service (FSIS) announced today. The RTE, frozen andouille sausages were produced on May 14, 2019. The following products are subject to recall: [View Labels (PDF Only)]

- Varying weights of vacuum-packed, individually-sealed packages containing "SAN GIUSEPPE SALAMI CO. BY GIACOMO ANDOUILLE SAUSAGE (SMOKE FLAVORING ADDED)" with a "Sell By: 11/10/2019" date on the case labels.

The products subject to recall bear establishment number "EST. 21556" inside the USDA mark of inspection. These items were shipped to retail and institutional locations in Greensboro, N.C. The firm contacted FSIS after they received a report that a consumer found a piece of a metal ring in the product.

There have been no confirmed reports of adverse reactions due to consumption of these products. FSIS has received no additional reports of injury or illness from consumption of these products. Anyone concerned about an injury or illness should contact a healthcare provider.

FSIS is concerned that some product may be in institutional and consumers' freezers. Consumers who have purchased these products are urged not to consume them and institutions should not serve these products. These products should be thrown away or returned to the place of purchase. FSIS routinely conducts recall effectiveness checks to verify recalling firms notify their customers of the recall and that steps are taken to make certain that the product is no longer available to consumers. When available, the retail distribution list(s) will be posted on the FSIS website at www.fsis.usda.gov/recalls.

Consumers and members of the media with questions about the recall can contact San Giuseppe Salami Co. by Giacomo's Plant Managers Francisco Grijalva or Giacomo Santomauro at (336) 586-7003. or at Francesco@sgsalami.com and Giacomo@sgsalami.com, respectively.

Consumers with food safety questions can "Ask Karen," the FSIS virtual representative available 24 hours a day at AskKaren.gov or via smartphone at m.askkaren.gov. The toll-free USDA Meat and Poultry Hotline 1-888-MPHotline (1-888-674-6854) is available in English and Spanish and can be reached from 10 a.m. to 6 p.m. (Eastern Time) Monday through Friday. Recorded food safety messages are available 24 hours a day. The online Electronic Consumer Complaint Monitoring System can be accessed 24 hours a day at: http://www.fsis.usda.gov/reportproblem.

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For full details on the recall, please visit: https://www.fsis.usda.gov/wps/portal/fsis/topics/recalls-and-public-health-alerts/recall-case-archive/archive/2019/recall-071-2019-release

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FEEDING AMERICA RECALL NOTICE

The following message is being sent to all authorized member recall personnel.

COMPANY ANNOUNCEMENT

Keurig Dr Pepper Announces Voluntary Withdrawal of Unflavored Peñafiel Mineral Spring Water that Does Not Meet FDA Bottled Water Quality Standards

When a company announces a recall, market withdrawal, or safety alert, the FDA posts the company's announcement as a public service. FDA does not endorse either the product or the company.

Read Announcement View Product Photos

Keurig Dr Pepper Announces Voluntary Withdrawal of Unflavored Peñafiel Mineral Spring Water that Does Not Meet FDA Bottled Water Quality Standards

Summary

Company Announcement Date:
June 21, 2019
FDA Publish Date:
June 21, 2019
Product Type:
Food & Beverages
Bottled Water
Reason for Announcement:
Recall Reason Description

Presence of Arsenic
Company Name:
Keurig Dr. Pepper
Brand Name:
Brand Name(s)
Penafiel
Product Description:
Product Description
Unflavored Mineral Spring Water

Company Announcement

BURLINGTON, MA and PLANO, TX - Keurig Dr Pepper today announced it will voluntarily withdraw Peñafiel unflavored mineral spring water products, imported from Mexico, due to the presence of violative levels of arsenic. Arsenic when present in the diet at very high levels, well above those detected in recent samples of Peñafiel, is associated with numerous chronic diseases. Water quality tests of Peñafiel samples conducted by an independent laboratory on behalf of Keurig Dr Pepper detected arsenic at levels that exceeded the FD!'s bottled water standards for mineral water of 10 ppb.

All unflavored Peñafiel mineral spring water products including 600mL and 1.5L of all date codes are included in this voluntary withdrawal. The product is packaged in PET bottle formats. Consumers who have this product in their possession can return it to their retailer for a full refund.

Peñafiel is a small brand in the U.S. and quantities in the marketplace are very limited, given that Keurig Dr Pepper has already begun to withdraw the products from the market. The Company has notified retailers that it will work with them to remove the product from the market.

Arsenic is found in nature, including in aquifers that are the source of mineral water and where levels can vary over time. Keurig Dr Pepper has recently installed enhanced filtration systems at its facilities that produce Peñafiel, and the product now being produced is well within regulatory guidelines.

No other Keurig Dr Pepper products are impacted by this voluntary removal. For further information, please contact the Keurig Dr Pepper Consumer Care hotline at (800) 696-5891 between the hours of 9:00 am and 8:00 pm EST, Monday through Friday. We are conducting this market action with the knowledge of the U.S. FDA.

About Keurig Dr Pepper

Keurig Dr Pepper (KDP) is a leading coffee and beverage company in North America, with annual revenue in excess of $11 billion. KDP holds leadership positions in soft drinks, specialty coffee and tea, water, juice and juice drinks and mixers, and markets the #1 single serve coffee brewing system in the U.S. The Company maintains an unrivaled distribution system that enables its portfolio of more than 125 owned, licensed and partner brands to be available nearly everywhere people shop and consume beverages. With a wide range of hot and cold beverages that meet virtually any consumer need, KDP key brands include Keurig®, Dr Pepper®, Green Mountain Coffee Roasters®, Canada Dry®, Snapple®, Bai®, Mott's®, CORE® and The Original Donut Shop®. The Company employs more than 25,000 employees and operates more than 120 offices, manufacturing plants, warehouses and distribution centers across North America. For more information, visitwww.keurigdrpepper.comExternal Link Disclaimer.

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* For additional local details, please contact the Health Department(s) for the area(s) your food bank serves.
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For full details on the recall, please visit: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/keurig-dr-pepper-announces-voluntary-withdrawal-unflavored-penafiel-mineral-spring-water-does-not?utm_campaign=Keurig%20Dr%20Pepper%20Announces%20Voluntary%20Withdrawal%20of%20Unflavored%20Pe%C3%B1afiel%20Mineral%20Spring%20Water&utm_medium=email&utm_source=Eloqua

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FEEDING AMERICA

RECALL NOTICE

The following message is being sent to all authorized member recall personnel.

COMPANY ANNOUNCEMENT

Perrigo Issues Voluntary Recall For Parent's Choice Advantage Infant Formula Milk-Based Powder With Iron

Read Announcement View Product Photos

Perrigo Issues Voluntary Recall For Parent's Choice Advantage Infant Formula Milk-Based Powder With Iron

Summary

Company Announcement Date:
June 21, 2019
FDA Publish Date:
June 21, 2019
Product Type:
Food & Beverages
Infant Formula & Foods
Reason for Announcement:
Recall Reason Description
Potential presence of metal foreign material
Company Name:
Perrigo Company plc
Brand Name:
Brand Name(s)
Parent's Choice
Product Description:
Product Description
Advantage Infant Formula Milk Based Powder with Iron

Company Announcement

Dublin, Ireland and Allegan, MI – Perrigo Company plc is issuing a voluntary nationwide recall of 35-ounce, 992-gram containers of Parent's Choice Advantage Infant Formula Milk-Based Powder with Iron. This product, sold exclusively at Walmart, is being recalled because of the

potential presence of metal foreign matter in a single lot of the product (C26EVFV). The total number of containers affected by this recall is 23,388.

No adverse events have been reported to date, and the recall is being initiated out of an abundance of caution stemming from a consumer report. No other products or retailers are affected by this recall.

Consumers who may have purchased the product should look for Lot Code C26EVFV with a "use by" date of February 26, 2021, which can be found on the bottom of the package. Any consumers who purchased the product should discontinue use and can visit any Walmart store for a refund.

Consumers with any health-related questions should contact their healthcare provider.

Consumers with questions about Parent's Choice Advantage Infant Formula Milk-Based Powder with Iron can contact Perrigo Consumer Affairs at 866-629-6181.

This recall is being conducted in consultation with the U.S. Food and Drug Administration (FDA).

About Perrigo
Perrigo Company plc is dedicated to making lives better by bringing high quality and affordable selfcare products that consumers trust everywhere they are sold. The Company is a leading provider of over-thecounter health and wellness solutions that enhance individual well-being by empowering consumers to proactively prevent or treat conditions that can be self-managed. Visit Perrigo online at http://www.perrigo.comExternal Link Disclaimer.

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* For additional local details, please contact the Health Department(s) for the area(s) your food bank serves.
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For full details on the recall, please visit: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/perrigo-issues-voluntary-recall-parents-choice-advantage-infant-formula-milk-

based-powder-iron?utm_campaign=Perrigo%20Issues%20Voluntary%20Recall%20For%20Parent%27s%20Choice%20Advantage%20Infant%20Formula%20Milk-Based%20Powder&utm_medium=email&utm_source=Eloqua