18-021 QA PRODUCT RECALL ORAJEL TEETHING PRODUCTS FROM CHURCH & DWIGHT TO BENZOCAINE INGREDIENT

Date: 15 JUN 2018 Recall: # 18-021

Scope: WORLDWIDE RETAIL FACILITIES

In consultation with the US Food & Drug Administration (FDA), Church & Dwight is voluntarily recalling Orajel teething products due to containing benzocaine.

Hazard: The use of benzocaine gels, sprays, ointments, solutions, and lozenges for mouth and gum pain, expecially in children under age 2, can lead to a serious—and sometimes fatal—condition called methemoglobinemia, in which the oxygen-carrying capacity of red blood cells is greatly reduced.

Incidents/Injuries: Out of an abundance of caution, Church & Dwight has decided to initiate the voluntary recall of this product.

Description: The following products are affected.

UPC	ITEM DESCRIPTION
310310319557	Orajel Instant Relief Daytime & Nighttime Gel
310310339401	Orajel Baby Nighttime Gel
310310033132	Orajel Baby Gel
310310340384	Orajel Baby Teething Swab

Remedy: Consumers should discontinue use of the item immediately and return to store for a full refund.

Consumer Contact: Consumers with questions may contact Church and Dwight Co. Inc's consumer relations at 1 (800) 833-9532, Monday through Friday 9:00 a.m. to 5:00 p.m., Eastern Standard Time.

Sold at: Worldwide Exchange facilities and other retailers.

Manufacturer(s): Church and Dwight Co.