

**Contact:**

Ilya Welfeld  
Seymour Public Relations  
804-431-3167  
news@sabra.com

**SABRA DIPPING COMPANY ISSUES VOLUNTARY RECALL OF CERTAIN HUMMUS PRODUCTS  
BECAUSE OF POSSIBLE HEALTH RISKS**

(November 19, 2016 Colonial Heights, VA)— Sabra Dipping Co., LLC is voluntarily recalling certain hummus products made prior to November 8, 2016 due to concerns over *Listeria monocytogenes*, which was identified at the manufacturing facility but not in tested finished product. The recall includes the products listed below; these were distributed to retail outlets, including food service accounts and supermarkets, in the U.S. and Canada.

*Listeria monocytogenes* is an organism, which can cause serious and sometimes fatal infections in young children, frail or elderly people, and others with weakened immune systems. Healthy individuals may suffer only short-term symptoms such as high fever, severe headaches, stiffness, nausea, abdominal pain and diarrhea. Listeria infection can cause miscarriages and stillbirths among pregnant women. The company is issuing this recall out of an abundance of caution.

Consumers with any product with a “Best Before” date up through January 23, 2017 are urged to discard it. Consumers can find code and “Best Before” date on the lid of each package.

**Costco Affected Item Numbers:**

Item Number	Description
818771	2-23.5 oz CLASSIC / GARLIC DUAL PK
32868	32 oz PINE NUT HUMMUS
851652	8 COUNT GRAB n GO HUMMUS + PRETZELS
464198	16 COUNT HUMMUS SINGLES
5170	5 LB CLASSIC HUMMUS TUB
32866	32 oz. ROASTED GARLIC HUMMUS

No other Sabra products are affected. In particular, Sabra products not included in the recall are: Sabra Organic Hummus, Sabra Salsa, Sabra Guacamole and Sabra Greek Yogurt Dips.

Consumers can contact Sabra Consumer Relations at 1-866-265-6761 for additional information from 9:00 am to 8:00 PM eastern time. For product reimbursement, consumers can contact [www.sabrahammusrecall.com](http://www.sabrahammusrecall.com). The company has subsequently taken steps to correct this matter.

The recall is being conducted with the knowledge of the U.S. Food and Drug Administration.