Golden State Medical Supply, Inc. Issues a Voluntary Nationwide Recall of Ibuprofen 800mg tablets. Due to potential aluminum particulate contamination localized on the surface of the tablets.

Company Contact:
Doug Blackburn, Vice President of Quality
(800) 284-8633, ext. 116

FOR IMMEDIATE RELEASE – November 23, 2026, Golden State Medical Supply, Incorporated (GSMS, Inc.) - Camarillo, CA has initiated a voluntary consumer level recall of Ibuprofen 800mg Tablets (NDC 51407-841-05) Lot# GS069303 due to potential aluminum particulate contamination localized on the surface of the tablets.

This voluntary recall only affects the product lot listed in the table below.

Product Description	NDC	Lot #	Exp. Date
IBUPROFEN 800MG TABLETS	51407-841-05	GS069303	01/2030

Continued use of the product from the affected lot may cause minor irritation and abrasion of the gastrointestinal track, e.g., stomach or intestinal linings. The likelihood of a related adverse event is remote given the size of the identified particulates (0.01 to 0.10 mm).

To date, GSMS is not aware of any adverse events associated with this lot.

Ibuprofen, a nonsteroidal anti-inflammatory drug (NSAID), is available in 800 mg tablets for oral administration. Ibuprofen tablets are indicated for relief of the signs and symptoms of rheumatoid arthritis and osteoarthritis, mild to moderate pain, and treatment of primary dysmenorrhea.

For more drug label information about Ibuprofen visit: DailyMed – Ibuprofen (nih.gov)

Consumers, patients, and caregivers are instructed to STOP the use of the medication, and immediately contact their physician, pharmacist, or healthcare provider. Any unused portion of the product should be returned to the pharmacy where it was obtained.

Consumers, patients, and caregivers are instructed to contact their physician or healthcare provider if they or the person they are giving care to has experienced any problems that may be related to taking or using this product.

The lot under GSMS, Inc.'s voluntary recall has been sold to McKesson. McKesson sites are instructed to immediately stop distribution, quarantine all remaining products in their control, and return the recalled product to GSMS, inc. They are also instructed to provide their customers, i.e., pharmacies and consumers, with a copy of GSMS, Inc.'s Recall Notification, Recall Response Form, and Letter to Consumers, Patients, and Caregivers.

Consumers with questions regarding this recall can contact GSMS, Inc, by:

- Phone: (800) 284-8633, ext. 116 (Monday through Friday between 7:30 AM to 4:00 PM, Pacific)
- Email: recalls@gsms.us

Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting program either online, by regular mail or by fax.

- Complete and submit the report Online: www.fda.gov/medwatch/report.htm
- Regular Mail or Fax: Download form www.fda.gov/MedWatch/getforms.htm or call 1-800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form, or submit by fax to 1-800-FDA-0178

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

Golden State Medical Supply, Inc. Product Labeling and Tablet Images:

Ibuprofen 800mg Tablets 51407-841-05



