

Mike DeWine
Governor**Jon Husted**
Lieutenant Governor**Dorothy Pelanda**
ODA Director**Amy Acton, MD, MPH**
ODH Director

DATE: November 13, 2019

TO: Health Commissioners, Directors of Environmental Health and Interested Parties

RE: Recall Announcement (ODA/ODH) 2019-186

Nature's Rx Issues Voluntary Nationwide Recall of Silver Bullet 10 Male Enhancement Capsules due to an Undeclared PDE-5 Inhibitor Found in The Product

Nature's Rx is voluntarily recalling lot: 01251ZX1, Expiry Date: 11/2022 of Silver Bullet (10 Male Enhancement Capsules). This recall has been initiated after an FDA laboratory analysis found the product to contain undeclared sildenafil, the active ingredient in Viagra, which is a PDE-5 inhibitor. The undeclared PDE-5 inhibitor in the product may pose serious health risks to consumers with underlying medical issues. For example, PDE-5 inhibitors may interact with nitrates found in some prescription drugs (such as nitroglycerin) lowering blood pressure to dangerous levels. Consumers with diabetes, high blood pressure, or heart disease often take nitrates.

To date, Nature's Rx has not received any reports of adverse events related to this recall.

The product, Silver Bullet (10 Male Enhancement Capsules), is used as a male enhancement nutritional supplement contained in blister foil sheets and packaged in small boxes containing 10 capsules with an expiry date: 11/2022 and lot number: 01251ZX1. Silver Bullet (10 Male Enhancement Capsules) was sold on an online website: www.naturalrx.net

Nature's Rx is notifying its customers by replacement or refund and is arranging for return or replacement of all recalled products. Consumers that have Silver Bullet (10 Male Enhancement Capsules), which is being recalled should stop using and return ship or discard the product.

Consumers with questions regarding this recall can contact Nature's Rx by phone number or e-mail address 5 days of the week from 10 AM to 6 PM PST. 1-888-925-5551, email us at: naturesrxsales@gmail.com or mail us at: 310 N Indian Hill Blvd Ste 600, Claremont, CA 91711

Consumers should contact their physician or healthcare provider if they have experienced any problems that may be related to taking or using this drug product.

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](#)
- Regular Mail or Fax: [Download form](#) 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.