



Ohio Department of Agriculture  
and  
Ohio Department of Health



**Governor** Ted Strickland  
**Lieutenant Governor** Lee Fisher

**ODA Director** Robert J. Boggs  
**ODH Director** Alvin D. Jackson, M.D.

To: Health Commissioners, Environmental Health Directors, Nursing Directors, and Other Interested Parties  
Subject: Recall Announcement (ODA/ODH) 2010-72  
RE: August 20, 2010

**Glow Industries, Inc. Issues Nationwide Recall of Mr. Magic Male Enhancer from Don Wands Amended**

Glow Industries, Inc., Perrysburg, OH, announced today that it is initiating a voluntary nationwide recall of the company's product sold under the name of Mr. Magic Male Enhancer from Don Wands. Glow Industries, Inc. is conducting this voluntary recall after being informed by the Food and Drug Administration (FDA) that lab analysis has found the Mr. Magic Male Enhancer from Don Wands capsules to contain Hydroxythiohomosildenafil and Sulfoildenafil, an analogue of Sildenafil, an FDA-approved drug used in the treatment of male Erectile Dysfunction (ED), making Mr. Magic Male Enhancer an unapproved new drug. These active ingredients are not listed on the product label. Product manufactured containing lot numbers 9041401, 251209 and 8121904 are included in this recall.

The Mr. Magic Male Enhancer recall includes:

Product Name	Lot Code	UPC Code
Mr. Magic 1 ct. Capsule Card	9041401 and 251209	648658123001
Mr. Magic 3 ct. Capsule Bottle	9041401 and 8121904	648658123018
Mr. Magic 6 ct. Capsule Bottle	9041401 and 8121904	648658123025
Mr. Magic 12 ct. Capsule Bottle	9041401 and 8121904	648658123032
Mr. Magic Display of 24; 1 ct. Capsule Cards	9041401 and 251209	648658123043
Mr. Magic 20 ct. Capsule Bottle	9041401 and 8121904	648658123056

The recall is being conducted as a precautionary measure. No illnesses or adverse effects have been reported to the company to date in connection with the product.

The undeclared ingredients may pose a threat to consumers because the analogue may interact with nitrates found in some prescription drugs (such as nitroglycerin) and may lower blood pressure to dangerous levels. Consumers with diabetes, high blood pressure, high cholesterol, or heart disease often take nitrates. ED is a common problem in men with conditions, and consumers may seek types of products to enhance sexual performance.

Glow Industries, Inc. advises any customer in possession of Mr. Magic Male Enhancer from Don Wands capsules to call (419) 350-2726, between the hours of 8 a.m and 5 p.m, Monday through Friday, Eastern Standard Time for instructions on the product return and credit process. Glow Industries, Inc. is notifying its distributors and retailers by a recall letter and phone calls to arrange for return of recalled product in their possession.

Any adverse events that may be related to the use of this product should be reported to the FDA's MedWatch Adverse Event Reporting program online [at [www.fda.gov/MedWatch/report.htm](http://www.fda.gov/MedWatch/report.htm)<sup>9</sup>], by returning the postage-paid FDA form 3500 [which may be downloaded from [www.fda.gov/MedWatch/getforms.htm](http://www.fda.gov/MedWatch/getforms.htm)<sup>10</sup>] by mail [to MedWatch, 5600 Fishers Lane, Rockville, MD 20852-9787] or fax [1-800-FDA-0178].

Customers with questions should contact Glow Industries, Inc. at (419) 350-2726.