

COMPANY ANNOUNCEMENT

Motto International Corp. Issues Voluntary Recall of Bull Platinum 30000, Stallion Platinum 30000, Rhino 7 Platinum 30000, and Panther Platinum 30000, Due to Presence of Undeclared Tadalafil

When a company announces a recall, market withdrawal, or safety alert, the FDA posts the company's announcement as a public service. FDA does not endorse either the product or the company.

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Summary

Company Announcement Date:

December 17, 2019

FDA Publish Date:

December 17, 2019

Product Type:

Drugs

Reason for Announcement:

Presence of Undeclared Tadalafil

Company Name:

Motto International Corp

Brand Name:

No Brand name Listed on Package

Product Description:

Bull Platinum 30000, Stallion Platinum 30000, Rhino 7 Platinum 30000, Panther Platinum 30000

Company Announcement


Motto International Corp. is voluntarily recalling all lots of Bull Platinum 30000, Stallion Platinum 30000, Rhino 7 Platinum 30000, and Panther Platinum 30000, to the consumer level. FDA analysis has found Bull Platinum 30000, Stallion Platinum 30000, Rhino 7 Platinum 30000, and Panther Platinum 30000 to be tainted with undeclared tadalafil. Tadalafil is an

FDA approved drug indicated for the treatment of male sexual enhancement. The presence of tadalafil in these products renders them unapproved drugs for which safety and efficacy has not been established and, therefore, subject to recall.



Consumption of a product with undeclared tadalafil may pose a risk to consumers who take prescription medications containing nitrates (such as nitroglycerin). The combination of tadalafil and nitrates may lower blood pressure to dangerous levels which can be life threatening. Consumers with diabetes, high blood pressure, high cholesterol, or heart disease often take nitrates and may be the population most likely to be affected. To date, Motto International Corp. has not received any reports of adverse events related to this recall.

The Bull Platinum 30000, Stallion Platinum 30000, Rhino 7 Platinum 30000, and Panther Platinum 30000 products are marketed as dietary supplements for male sexual enhancement and are packaged in 2, 4, and 10 capsule blister pack cards. The product can be identified by the product labels attached below. Product was sold locally to 2 Dallas convenience stores.

Motto International Corp. is notifying its customers by email and is arranging for replacement of recalled products, at the request of the customer. Consumers who purchased the product in October, 2019 should stop consuming it and return any unused product to Motto International Corp., 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.

Consumers with questions regarding this recall can contact Motto International Corp. via email at mottoic@gmail.com (<mailto:mottoic@gmail.com>) at any time or by phone at 469-644-4398  on Monday through Friday between the hours of 9 a.m. to 5 p.m. Central Time.

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 (</safety/medwatch-fda-safety-information-and-adverse-event-reporting-program/reporting-serious-problems-fda>)
- Regular Mail or Fax: Download form (</safety/medical-product-safety-information/medwatch-forms-fda-safety-reporting>) 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.

Company Contact Information

Consumers:

Sam Lee

☎ 469-644-4398

✉ mottoic@gmail.com (mailto:mottoic@gmail.com)

Product Photos









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