Company Announcement

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.

Regeneca Worldwide, a Division of Vivaceuticals, Inc., Issues a Nationwide Recall of Its Products

For Immediate Release

March 9, 2017

Contact

Consumers

Regeneca Support
support@regeneca.com (mailto:support@regeneca.com)

Announcement

Regeneca Worldwide, a division of VivaCeuticals, Inc. is conducting a nationwide recall of its entire line of herbal and dietary supplement products pursuant to a Consent Decree entered by the federal court for the Central District of California. This recall applies to all products manufactured and sold between June 1, 2011 and February 8, 2017. These products include, but are not limited to RegeneSlim, RegenErect, RegeneArouse, RegeneBlend, RegeneBoost, RegeneBlast, and RegeneFit. ALL LOT #s ARE INCLUDED IN THIS RECALL. Regeneca products were sold nationwide online using the company's websites, and through a direct sales force within the United States and Puerto Rico for both consumption and retail sales.

The company has presently ceased doing business in the United States.

Consumers who have purchased Regeneca products subject to this recall are advised to immediately stop using the product and are urged to return it to the place of purchase for a possible exchange. Consumers with questions may contact the company via email to support@regeneca.com (mailto:support@regeneca.com).

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 (http://www.fda.gov/medwatch/report.htm)
- **Regular Mail or Fax**: Download form www.fda.gov/MedWatch/getforms.htm (http://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

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