Humana.

Drug recall notice for EpiPen® and EpiPen Jr® Auto-Injector

The maker of EpiPen® and EpiPen Jr® Auto-Injectors has voluntarily recalled select lots distributed in the U.S. and other markets. Lot numbers are assigned to products by manufacturers and can be used as a reference in the event of a recall. This recall was initiated due to reports of possible device malfunctions that could result in the devices failing to activate or requiring increased force to activate in an emergency or life-threatening situation.

This recall is intended for the following products:

National drug code (NDC)	Product	Lot numbers	Expiration
49502-501-02 49502-500-02	EpiPen Jr 2-Pak® Auto-Injector, 0.15 mg EpiPen 2-Pak® Auto-Injector, 0.3 mg	5GN767; 5GN773; 5GM631; 5GM640; 6GN215; 6GM082; 6GM072; 6GM081; 6GM088; 6GM199; 6GM091; 6GM198; 6GM087	April 2017 through October 2017

What this means for you

- Check the lot number on your carton or device to see if the EpiPen® and EpiPen Jr® Auto-Injector is affected by the recall.
- If your EpiPen® and EpiPen Jr® Auto-Injector has been recalled, contact Stericycle at **1-877-650-3494** to obtain a voucher code for your free replacement product. Stericycle will also provide you with a pre-paid return package to ship the product back to Stericycle. Stericycle's hours of operation are Monday through Friday, 8 a.m. to 10 p.m. ET, and Saturday and Sunday 8 a.m. to 5 p.m. ET.
- Visit your pharmacy with your voucher information to redeem your free replacement.
- Send your recalled product to Stericycle. Do not return any devices affected by the recall until you have your replacement in hand.
- If you have questions regarding this recall, please contact Mylan Customer Relations at **1-800-796-9526**. Representatives are available Monday through Friday from 8 a.m. to 8 p.m. ET or by email at **customer.service@mylan.com**.
- **It is important that you continue to carry your current EpiPen® Auto-Injector until you receive a replacement device.**
- 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.
    - **Online**: Complete and submit the report: **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**.

If you have any questions, please talk to your physician or healthcare professional. You may also call the number on the back of your Humana member ID card. Our automated phone system may answer your call on Saturdays, Sundays, and some public holidays. Please leave your name and telephone number and we'll call you back by the end of the next business day. For 24-hour service you can visit **Humana.com**.