NuFACTOR Specialty Pharmacy

Dear Valued Patient,

NuFACTOR was recently notified that the U.S. Food and Drug Administration is alerting consumers to Meridian Medical Technologies' **voluntary recall of 13 lots of Mylan's EpiPen and EpiPen Jr (epinephrine injection) Auto-Injector** products used for emergency treatment of severe allergic reactions.

This recall is due to the potential that these devices may contain a defective part that may result in the devices failure to activate.

EpiPen Return and Replacement Process

If you think you may be impacted by this recall, it is very important to **contact Stericycle at (877) 650-3494**. Stericycle will verify if your EpiPen 2-Pak® or EpiPen Jr 2-Pak® cartons are from one of the recalled lots.

- **Prior to calling Stericycle**, confirm if you are in possession of a recalled EpiPen product by checking if the lot number matches any of the lot numbers listed in the table below. If so, you need to contact Stericycle at (877) 650-3494. If not, your EpiPen product is not affected by the recall and there is no further action necessary.

- **If you are in possession of a recalled EpiPen product**, Stericycle will initiate the process of providing a container to return the recalled medication and issue a voucher to redeem a free replacement from your pharmacy. It is important that patients continue to carry their current EpiPen Auto-Injector until they receive a replacement device. Patients should not return any devices affected by the recall until they have received their voucher.

    - Stericycle also will collect your contact information and will begin contacting patients back on Monday, April 3, with voucher information to redeem a free replacement product.
    - Patients may receive either EpiPen Auto-Injector or Mylan's authorized generic for EpiPen Auto-Injector at the pharmacy as a replacement based on availability. The authorized generic has the exact same drug formulation, has the exact same operating instructions and is therapeutically equivalent to EpiPen Auto-Injector, and may be substituted for EpiPen Auto-Injector.

*****Please refer to the following page for affected lots. *****

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Affected Lots Include:

Product/Dosage	NDC Number on Carton	Lot Number	Expiration Date
EpiPen Jr 2-Pak® Auto-Injectors, 0.15 mg	49502-501-02	5GN767	April 2017
EpiPen Jr 2-Pak® Auto-Injectors, 0.15 mg	49502-501-02	5GN773	April 2017
EpiPen 2-Pak® Auto-Injectors, 0.3 mg	49502-500-02	5GM631	April 2017
EpiPen 2-Pak® Auto-Injectors, 0.3 mg	49502-500-02	5GM640	May 2017
EpiPen Jr 2-Pak® Auto-Injectors, 0.15 mg	49502-501-02	6GN215	September 2017
EpiPen 2-Pak® Auto-Injectors, 0.3 mg	49502-500-02	6GM082	September 2017
EpiPen 2-Pak® Auto-Injectors, 0.3 mg	49502-500-02	6GM072	September 2017
EpiPen 2-Pak® Auto-Injectors, 0.3 mg	49502-500-02	6GM081	September 2017
EpiPen 2-Pak® Auto-Injectors, 0.3 mg	49502-500-02	6GM088	October 2017
EpiPen 2-Pak® Auto-Injectors, 0.3 mg	49502-500-02	6GM199	October 2017
EpiPen 2-Pak® Auto-Injectors, 0.3 mg	49502-500-02	6GM091	October 2017
EpiPen 2-Pak® Auto-Injectors, 0.3 mg	49502-500-02	6GM198	October 2017
EpiPen 2-Pak® Auto-Injectors, 0.3 mg	49502-500-02	6GM087	October 2017

Please be aware that Mylan has informed NuFACTOR that we cannot provide you with any assistance. Patients must manage the return and replacement process of recalled EpiPens on their own.

To learn more regarding this recall, please visit - www.mylan.com/en/epipenrecall or www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm550170.htm.

Sincerely,

NuFACTOR Specialty Pharmacy
(800) 323-6832