EpiPen (epinephrine injection, USP) and EpiPen Jr® (epinephrine injection, USP) Auto-Injectors

On 03/31/2017, Mylan recalled lot numbers (5GN767, 5GN773, 5GM631, 5GM640, 6GN215, 6GM082, 6GM072, 6GM081, 6GM088, 6GM199, 6GM091, 6GM198, 6GM087) of EpiPen (epinephrine injection, USP) and EpiPen Jr® (epinephrine injection, USP) Auto-Injectors (NDC# 49502-501-02, 49502-500-02) for failure to activate the device due to a potential defect. The U.S. Food and Drug Administration (FDA) has issued a Class I of the affected medications. More information about the recall is at: http://www.fda.gov/Safety/Recalls/EnforcementReports/default.htm.

The detailed information of the recalled products is listed below:

Recalled Drug: EpiPen (epinephrine injection, USP) and EpiPen Jr® (epinephrine injection, USP) Auto-Injectors
NDC Number: 49502-501-02, 49502-500-02
Lot Numbers: 5GN767, 5GN773, 5GM631, 5GM640, 6GN215, 6GM082, 6GM072, 6GM081, 6GM088, 6GM199, 6GM091, 6GM198, 6GM087
Expiration Date: Apr-17, May-17, Sep-17, Oct-17

What you should do:

- **Do not continue to use EpiPen (epinephrine injection, USP) and EpiPen Jr® (epinephrine injection, USP) Auto-Injectors if it has been recalled**
- Check your prescription label to see if you have EpiPen (epinephrine injection, USP) and EpiPen Jr® (epinephrine injection, USP) Auto-Injectors.
- If you have the medication, contact the pharmacy that you received it from.
- Adverse events from the use of prescription drugs should be reported to FDA by visiting www.fda.gov/medwatch/report.htm or calling 1-800-FDA-0178.

Oxycodone Hydrochloride Oral Solution, USP 5 mg/5 mL

On 01/18/2017, VistaPharm recalled (Lot Numbers: 427900, 426700, 424800, 423600, 420800, 416300, 407700, 407300, 405900, 403900, 426900, 404700, 390200) of Oxycodone Hydrochloride Oral Solution, USP 5 mg/5 mL (NDC# 66689-401-50, 66689-403-16), the purified water used to manufacture the drug products may have been contaminated with Burkholderia cepacia. The U.S. Food and Drug Administration (FDA) has issued a Class II of the affected medications. More information about the recall is at: http://www.fda.gov/Safety/Recalls/EnforcementReports/default.htm.
The detailed information of the recalled product is listed below:

Recalled Drug: Oxycodone Hydrochloride Oral Solution, USP 5 mg/5 mL
NDC Number: 66689-401-50, 66689-403-16
Lot Numbers: 427900, 426700, 424800, 423600, 420800, 416300, 407700, 407300, 405900, 403900, 426900, 404700, 390200
Expiration Date: 11/17, 10/17, 09/17, 08/17, 06/17, 05/17, 02/17

What you should do:

- ☐ **Do not continue to use Oxycodone Hydrochloride Oral Solution, USP 5 mg/5 mL if it has been recalled**
- ☐ Check your prescription label to see if you have Oxycodone Hydrochloride Oral Solution, USP 5 mg/5 mL.
- ☐ If you have the medication, contact the pharmacy that you received it from.
- ☐ Adverse events from the use of prescription drugs should be reported to FDA by visiting www.fda.gov/medwatch/report.htm or calling 1-800-FDA-0178.

GlucaGen HypoKit (glucagon [rDNA origin] for injection).

On 09/08/2016, Novo Nordisk recalled (FS6X270, FS6X296, FS6X538, FS6X597, FS6X797, FS6X875) of GlucaGen HypoKit (glucagon [rDNA origin] for injection) (NDC# 0169-7065-15) Defective delivery system: detached needles on the syringe in the kit.

The U.S. Food and Drug Administration (FDA) has issued a Class I of the affected medications. More information about the recall is at: http://www.fda.gov/Safety/Recalls/EnforcementReports/default.htm.

The detailed information of the recalled product is listed below:

Recalled Drug: GlucaGen HypoKit (glucagon [rDNA origin] for injection).
NDC Number: 0169-7065-15
Lot Numbers: FS6X270, FS6X296, FS6X538, FS6X597, FS6X797, FS6X875
Expiration Date: 9/30/2017

What you should do:

- ☐ **Do not continue to use GlucaGen HypoKit (glucagon [rDNA origin] for injection) if it has been recalled.**
- ☐ Check your prescription label to see if you have GlucaGen HypoKit (glucagon [rDNA origin] for injection).
- ☐ If you have the medication, contact the pharmacy that you received it from.
- ☐ Adverse events from the use of prescription drugs should be reported to FDA by visiting www.fda.gov/medwatch/report.htm or calling 1-800-FDA-0178.

Senna Syrup (sennosides) 8.8 mg, 8 fl. oz. (237 mL) bottle

On 8/8/2016, Major Pharmaceuticals recalled Senna Syrup (sennosides) Syrup (NDC# 0904-6289-09). This recall was initiated as a precautionary measure due to a potential risk of product contamination with the bacteria B. cepacia. The U.S. Food and Drug Administration (FDA) has issued a Class II of the affected medications. More information about the recall is at: http://www.fda.gov/Safety/Recalls/EnforcementReports/default.htm.
The detailed information of the recalled product is listed below:

Recalled Drug: Senna Syrup (sennosides) 8.8 mg
NDC Number: 0904-6289-09
Lot Number: 20391517, 20391518, 20391519, 20391601, 20391602, 20391604, 20391605, 20391608.
Expiration Date: 09/17, 10/17, 11/17, 01/18, 02/18, 03/18, 06/18.

What you should do:

- ☐ **Do not continue to use Senna Syrup (sennosides) if it has been recalled**
- ☐ Check your prescription label to see if you have Senna Syrup (sennosides).
- ☐ If you have the medication, contact the pharmacy that you received it from.
- ☐ Adverse events from the use of prescription drugs should be reported to FDA by visiting www.fda.gov/medwatch/report.htm or calling 1-800-FDA-0178.

Senexon Liquid (sennosides) 8.8 mg, 8 fl oz. (237 mL)

On 8/8/2016, Rugby Laboratories recalled Senexon Liquid (sennosides) (NDC# 0536-1000-59). This recall was initiated as a precautionary measure due to a potential risk of product contamination with the bacteria B. cepacia. The U.S. Food and Drug Administration (FDA) has issued a Class II of the affected medications. More information about the recall is at: http://www.fda.gov/Safety/Recalls/EnforcementReports/default.htm.

The detailed information of the recalled product is listed below:

Recalled Drug: Senexon Liquid (sennosides) 8.8 mg
NDC Number: 0536-1000-59

What you should do:

- ☐ **Do not continue to use Senexon Liquid (sennosides) 8.8 mg if it has been recalled**
- ☐ Check your prescription label to see if you have < Senexon Liquid (sennosides) 8.8 mg.
- ☐ If you have the medication, contact the pharmacy that you received it from.
- ☐ Adverse events from the use of prescription drugs should be reported to FDA by visiting www.fda.gov/medwatch/report.htm or calling 1-800-FDA-0178.

Diocto Syrup (docusate sodium), 60 mg/15 mL

On 8/8/2016, Rugby Laboratories recalled Diocto Syrup (docusate sodium), 60 mg/15 mL (NDC# 0536-1001-85). This recall was initiated as a precautionary measure due to a potential risk of product contamination with the bacteria B. cepacia. The U.S. Food and Drug Administration (FDA) has issued a Class II of the affected medications. More information about the recall is at: http://www.fda.gov/Safety/Recalls/EnforcementReports/default.htm.
The detailed information of the recalled product is listed below:

Recalled Drug: Diocto Syrup (docusate sodium), 60 mg/15 mL
NDC Number: 0536-1001-85

What you should do:

- ☐ **Do not continue to use Diocto Syrup (docusate sodium), 60 mg/15 mL if it has been recalled**
- ☐ Check your prescription label to see if you have Diocto Syrup (docusate sodium), 60 mg/15 mL.
- ☐ If you have the medication, contact the pharmacy that you received it from.
- ☐ Adverse events from the use of prescription drugs should be reported to FDA by visiting www.fda.gov/medwatch/report.htm or calling 1-800-FDA-0178.

Sennazon (sennosides) Syrup, 8.8 mg, 8 fl. oz. (237 mL) bottle

On 8/8/2016, Bayshore Pharmaceuticals recalled Sennazon (sennosides) Syrup (NDC# 76518-100-08). This recall was initiated as a precautionary measure due to a potential risk of product contamination with the bacteria B. cepacia. The U.S. Food and Drug Administration (FDA) has issued a Class II of the affected medications. More information about the recall is at: http://www.fda.gov/Safety/Recalls/EnforcementReports/default.htm.
The detailed information of the recalled product is listed below:

Recalled Drug: Sennazon (sennosides) Syrup
NDC Number: 76518-100-08

What you should do:

- ☐ **Do not continue to use Sennazon (sennosides) Syrup if it has been recalled**
- ☐ Check your prescription label to see if you have Sennazon (sennosides) Syrup.
- ☐ If you have the medication, contact the pharmacy that you received it from.
- ☐ Adverse events from the use of prescription drugs should be reported to FDA by visiting www.fda.gov/medwatch/report.htm or calling 1-800-FDA-0178.

Albuterol Sulfate Syrup, 2 mg/ 5 mL, 473 mL bottle

On 06/22/2016, Teva Pharmaceuticals recalled Lot # 95113 of Albuterol Sulfate Syrup, 2 mg/ 5 mL, 473 mL bottle (NDC# 0093-0661-16) due to the presence of foreign substances; presence of black particles described generically as cellulose-based bundles of brown fibrous material. The U.S. Food and Drug Administration (FDA) has issued a Recall Class II of the affected medications. More information about the recall can be found at: http://www.fda.gov/Safety/Recalls/EnforcementReports/default.htm.

The detailed information of the recalled product is listed below:

Recalled Drug: Albuterol Sulfate Syrup, 2 mg/ 5 mL
NDC Number: 0093-0661-16
Lot Numbers: 95113
Expiration Date: 01/2017

What you should do:

- **Do not continue to use Albuterol Sulfate Syrup, 2 mg/ 5 mL if it has been recalled**
- Check your prescription label to see if you have Albuterol Sulfate Syrup, 2 mg/ 5 mL.
- If you have the medication, contact the pharmacy that you received it from.
- Adverse events from the use of prescription drugs should be reported to FDA by visiting www.fda.gov/medwatch/report.htm or calling 1-800-FDA-0178.