Sun Pharma – Recall of vecuronium

- On February 4, 2019, Sun Pharma announced a voluntary, consumer-level recall of several lots of vecuronium injection due to foreign matter identified as glass detected in one vial to date.
- The recalled lots were distributed January 12, 2018.

Product Description	NDC#	Lot# (Expiration Date)
Vecuronium bromide for injection 20 mg (10 x 20 mg vials, lyophilized powder)	47335-932-44; 47335-932-40	JKS0400A (3/2019)
Vecuronium bromide for injection 10 mg (10 x 10 mg vials, lyophilized powder)	47335-931-44; 47335-931-40	JKS0443A (3/2019); JKS0444A (3/2019); JKS0477A (3/2019)

- Vecuronium bromide is indicated as an adjunct to general anesthesia, to facilitate endotracheal intubation and to provide skeletal muscle relaxation during surgery or mechanical ventilation.
- Per Sun Pharma, use of the recalled product can possibly pose a risk to patient safety.
- Anyone with an existing inventory of the recalled product should stop use and distribution, and quarantine the product immediately.
- For more information regarding this recall, contact Inmar (appointed company for Sun Pharma) at **1-800-967-5952**.