



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**.



OptumRx® specializes in the delivery, clinical management and affordability of prescription medications and consumer health products. We are an Optum® company — a leading provider of integrated health services. Learn more at optum.com.

All Optum® trademarks and logos are owned by Optum, Inc. All other brand or product names are trademarks or registered marks of their respective owners.

This document contains information that is considered proprietary to OptumRx and should not be reproduced without the express written consent of OptumRx.

RxNews® is published by the OptumRx Clinical Services Department.

©2019 Optum, Inc. All rights reserved.