

Drug Recall

DATE OF RECALL: October 3, 2023

DRUG NAME: Various products

RECALLING FIRM: Hospira, Inc.

REASON FOR RECALL: This recall was issued due to the potential for presence of glass particulate matter.

FDA LINK: N/A

OTHER DETAILS: N/A

RECALLED PRODUCT:

| Product Description | NDC Number | Lot Number | Expiration Date |
|--|---------------|------------|-----------------|
| 4.2% Sodium Bicarbonate Injection, USP Glass ABBOJECT® Syringe Carton | 00409-5534-24 | GJ5007 | 08/01/2024 |
| 4.2% Sodium Bicarbonate Injection, USP Glass ABBOJECT® Syringe Case | 00409-5534-14 | GJ5007 | 08/01/2024 |
| 1% Lidocaine HCl Injection, USP LIFESHIELD® Glass ABBOJECT® Syringe Carton | 00409-4904-11 | 42290DK | 06/01/2024 |
| 1% Lidocaine HCl Injection, USP LIFESHIELD® Glass ABBOJECT® Syringe Case | 00409-4904-34 | 42290DK | 06/01/2024 |
| 2% Lidocaine HCl Injection, USP LIFESHIELD® Glass ABBOJECT® Syringe Carton | 00409-4903-11 | GH6567 | 07/01/2024 |
| 2% Lidocaine HCl Injection, USP LIFESHIELD® Glass ABBOJECT® Syringe Case | 00409-4903-34 | GH6567 | 07/01/2024 |