



CALIFORNIA DRUG RECALL INFORMATION SHEET

Hospira, Inc. Issues a Voluntary Nationwide Recall for 4.2% Sodium Bicarbonate Injection, USP and 1% and 2% Lidocaine HCl Injection, USP Due to the Potential for Presence of Glass Particulate Matter

Recall Date	Product Description	Recalling Firm	Recall Reason
10/02/2023	4.2% Sodium Bicarbonate Injection, USP, 5 mEq/10mL vial 1% Lidocaine HCl Injection, USP, 50 mg/5mL vial 2% Lidocaine HCl Injection, USP, 100 mg/5mL vial	Hospira, Inc., a Pfizer company	Potential for presence of glass particulate matter Potential complications related to injection of visible and subvisible inert particles include inflammation of a vein, granuloma, and blockage of blood vessels or life-threatening blood clot events. The frequency and severity of these adverse events could vary depending upon a variety of factors including the size and number of particles in the drug product, patient comorbidities (such as age, compromised organ function), and presence or absence of vascular anomalies.

Recall Class	Product Identification	Distribution	Affected Dates
N/A	<p>4.2% Sodium Bicarbonate Injection, USP, Glass ABBOJECT® Syringe</p> <p>5 mEq/10mL, (0.5 mEq/mL) 1 vial and injector/ carton 10 cartons/ bundle Case pack 5 X 10- 10mL</p> <p>NDC #: Carton 0409-5534-24 Case 0409-5534-14 Lot Number: 42290DK</p>	Nationwide	Product Expiration Date: 1 August 2024
N/A	<p>1% Lidocaine HCl Injection, USP, LIFESHIELD® Glass ABBOJECT® Syringe</p> <p>50 mg/5mL (10 mg/mL) 1 vial and injector/ carton 10 cartons/ bundle Case pack 5 X 10- 5mL</p> <p>NDC #: Carton 0409-4904-11 Case 0409-4904-34 Lot Number: 42290DK</p>	Nationwide	Product Expiration Date: 1 June 2024
N/A	<p>2% Lidocaine HCl Injection, USP LIFESHIELD, Glass ABBOJECT Syringe</p> <p>100 mg/5mL (20 mg/mL) 1 vial and injector/ carton 10 cartons/ bundle Case pack 5 X 10- 5mL</p> <p>NDC #: Carton 0409-4903-11 Case 0409-4903-34 Lot Number: GH6567</p>	Nationwide	Product Expiration Date: 1 July 2024

FOR ADDITIONAL INFORMATION, PLEASE VISIT [THE FDA WEBSITE](#)

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