

The Board of Pharmacy has received notice of the following product recall. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

Par Health USA, LLC ("Par Health") has identified the potential for the presence of particulate matter identified as cellulose in the product solution for lot 88802 and identified as stopper material in the product solution for lot 87558 in Dexmedetomidine HCl in 0.9% Sodium Chloride Injection (400 mcg/100 ml) 10-pack carton.

This recall is being conducted to the **hospital/clinic level**. All locations which have received direct shipment of these lots from Par Health have been sent a recall notification packet.

Shipments from these lots occurred between 05/13/2025 and 06/29/2025.

Product	Distributed By	NDC Number	Lot #	Exp. Date
Dexmedetomidine HCl in 0.9% Sodium Chloride Injection (400 mcg/100 mL) 10-pack carton	Par Pharmaceutical*	42023-187-10 (carton)	88802	04/2027
		42023-187-01 (vial)		
Dexmedetomidine HCl in 0.9% Sodium Chloride Injection (400 mcg/100 mL) 10-pack carton	Par Pharmaceutical*	42023-187-10 (carton)	87558	02/2027
		42023-187-01 (vial)		

*Par Pharmaceutical, Inc. is a predecessor company to Par Health USA, LLC, the current distributor of the product.