

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.

This is to communicate that Par Health is conducting a drug recall of the following lots of Brevital® Sodium 500 mg/vial CIV (Methohexital Sodium for Injection, USP).

Par Health USA, LLC ("Par Health") has identified the potential for the presence of particulate matter identified as cellulose in the product solution.

This recall is being conducted to the hospital/clinic level.

Product	Distributed By	NDC Number	Lot Number	Exp. Date
Brevital® Sodium 500 mg/vial CIV (Methohexital Sodium for Injection, USP) Unit Carton	Par Pharmaceutical*	42023-105-01	86930	02/2028
Brevital® Sodium 500 mg/vial CIV (Methohexital Sodium for Injection, USP) Unit Carton	Par Pharmaceutical*	42023-105-01	88623	04/2028