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 inform you that Chiesi USA, Inc. (“Chiesi”), is initiating a recall of two (2) lot numbers of CLEVIPREX- clevipidine emulsion, NDC 10122-611-10 (CLEVIPREX 50MG/100ML Vial), from the wholesale distribution channels and direct customers ONLY. The lots subject to the recall were sold from June 1, 2026 to June 30, 2026, and are listed below.

This action follows a deviation in manufacturing where there is a possible detachment or loosening of the crimped metal ring together with the flip-off cap during opening of Cleviprex vials. The product met all specification requirements at release, but we are initiating this recall as an additional precautionary measure.

Product Description	NDC	Lot Number	Expiration Date
CLEVIPREX 50MG/100ML Vial	10122-611-10	1235921	10/2027
CLEVIPREX 50MG/100ML Vial	10122-611-10	1237454	10/2027