The Board of Pharmacy has received notice of the following product recall:

Product Name: Acetaminophen Injection 1000mg/100mL (10mg/mL)
Lot #/Expiration Date:
NDC: 55150-307-24 – Outer Package; 55150-307-01 – Single Vial
Batch Number/Expiration Date:
CAT200002-09/2022; CAT200004-09/2022; CAT200005-09/2022; CAT200008-09/2022; CAT200009-09/2022; CAT200013-10/2022; CAT200014-10/2022; CAT200015-10/2022; CAT200016-10/2022; CAT200017-10/2022; and CAT200018-10/2022.

AuroMedics Pharma LLC has initiated a voluntary recall to the retail level of Acetaminophen Injection 1000mg/100mL distributed in the USA due to observed discoloration of the product. The package insert for this product states that the solution is clear and colorless. Some vials have been observed to be pale yellow to yellow which does not meet the approved product description. Additionally, some vials exhibiting discoloration have been found to be out of specification for certain parameters including color absorbance and pH. This recall is at the retail level.

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