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

Description: Ketorolac Tromethamine Injection, USP 30mg/mL
NDC: 0641-6042-25
Lot/Expiry Dates:
038366 / Mar-2020
048365 / Apr-2020
048367 / Apr-2020
078301 / Jul-2020
078303 / Jul-2020
118358 / Nov-2020
019413 / Jan-2021
029353 / Feb-2021
038368 / Mar-2020

Description: Ketorolac Tromethamine Injection, USP 30mg/mL
NDC: 0641-6043-25
Lot/Expiry Dates:
058314 / May-2020
078305 / Jul-2020
078307/ Jul-2020
118362 / Nov-2020

The decision to recall the product lots is in response to small black particles being noted during routine retain visual inspection of Ketorolac lots.

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.