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.

Product name/size	**NDC Number**	**Product Code**	**Batch Number**	**Exp. Date**	**First Ship Date**	**Last Ship Date**
Ketorolac Tromethamine Injection, USP, 60 mg / 2 mL (30 mg / mL), 2 mL fill in a 2 mL amber vial	63323-162-02	160202	6121125	02/2021	4/10/2019	5/23/2019
Ketorolac Tromethamine Injection, USP, 30 mg / 2 mL (30 mg / mL), 1 mL fill in a 2 mL amber vial	**63323-162-01**	**160201**	**6121083**	**02/2021**	**3/28/2019**	**9/3/2019**

Fresenius Kabi USA LLC is recalling the batches above to the user level due to particulate matter found in reserve sample vials.

NOTE: This notice contains one batch noted above in bold. A previous recall dated 12/17/2020 did not include product code 160201, batch number 6121083. All other information remains unchanged.