



Drug Recall

DATE OF RECALL: June 17, 2020

DRUG NAME: Ketorolac Tromethamine 30mg/ml Injection

RECALLING FIRM: Fresenius Kabi USA, LLC

REASON FOR RECALL: This recall was issued due to the presence of particulate matter found in reserve sample vials at the firm.

FDA LINK: <https://www.accessdata.fda.gov/scripts/ires/index.cfm?Product=181109>

OTHER DETAILS: See FDA link.

RECALLED PRODUCT:

Product Description	NDC Number	Affected Lots	Expiration Date
Ketorolac Tromethamine 30mg/ml Injection, Single Dose 1ml Vials	63323-0162-01	6118737	04/2020
		6118902	04/2020
		6119052	05/2020
		6119752	08/2020
		6122349	07/2021
		6122538	09/2021
Ketorolac Tromethamine 30mg/ml Injection, Single Dose 1ml Vials	63323-0162-00	6118737	04/2020
		6118902	04/2020
		6119052	05/2020
		6119752	08/2020
		6122349	07/2021
		6122538	09/2021