Mass General Brigham
Health Plan

Drug Recall

DATE OF RECALL: November 1, 2023

DRUG NAME: LEADER Brand Ophthalmic Sterile Drops supplied

RECALLING FIRM: Velocity Pharma, LLC and distributed by Cardinal Health, Inc.

REASON FOR RECALL: This recall was issued due to unsanitary conditions in the manufacturing facility and positive bacterial test results from environmental sampling of critical drug production areas in the facility.

FDA LINK: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/cardinal-health-inc-issues-voluntary-nationwide-recall-certain-leadertm-brand-eye-drops-supplied

OTHER DETAILS: N/A

RECALLED PRODUCT:

Product Description	NDC Number	Lot Number	Expiration Date
Eye Irritation Relief (Polyvinyl Alcohol, 0.5%, Povidone, 0.6%, and Tetrahydrozoline Hydrochloride, 0.05%) 0.5 FL OZ bottle (15 mL)	70000-0087-01	All	
Dry Eye Relief (Carboxymethylcellulose Sodium, 1%) 0.5 FL OZ bottle (15 mL)	70000-0089-01	All	
Lubricant Eye Drops (Carboxymethylcellulose Sodium, 0.5%) 0.5 FL OZ bottle (15 mL)	70000-0090-01	All	
Lubricant Eye Drops (Carboxymethylcellulose Sodium, 0.5%) 2 bottles, 0.5 FL OZ (15 mL) each	70000-0090-02 (carton) 70000-0090-01 (bottle)	All	
Dry Eye Relief (Polyethylene Glycol 400,0.4% and Propylene Glycol, 0.3%) 0.33 FL OZ bottle (10 mL)	70000-0088-01	All	
Lubricant Eye Drops (Propylene Glycol, 0.6%) 0.33 FL OZ bottle (10 mL)	70000-0587-01	All	