Mass General Brigham
Health Plan

Drug Recall

DATE OF RECALL: July 24, 2023

DRUG NAME: Famotidine 20mg Tablets

RECALLING FIRM: Glenmark Therapeutics, Inc.

REASON FOR RECALL: This recall was issued due to labeling: label error on declared strength: some cartons labeled and containing 20 mg may have an external label placed on the side of the carton indicating strength as 10 mg.

FDA LINK: N/A

OTHER DETAILS: N/A

RECALLED PRODUCT:

Product Description	NDC Number	Lot Number	Expiration Date
Famotidine 20mg Tablets	72657-0113-20	FA2022001B	03/2025