Mass General Brigham
Health Plan

Drug Recall

DATE OF RECALL: March 22, 2022

DRUG NAME: Accuretic (quinapril HCl/hydrochlorothiazide), Quinapril and Hydrochlorothiazide Tablets, and Quinapril HCl/Hydrochlorothiazide Tablets

RECALLING FIRM: Pfizer

REASON FOR RECALL: This recall was issued due to the presence of a nitrosamine, N-nitroso-quinapril, above the Acceptable Daily Intake (ADI) level.

FDA LINK: Pfizer Voluntary Nationwide Recall of Lots of ACCURETICTM (Quinapril HCl/Hydrochlorothiazide), Quinapril and Hydrochlorothiazide Tablets, and Quinapril HCl/Hydrochlorothiazide Tablets Due to N-NitrosoQuinapril Content | FDA

OTHER DETAILS: See FDA Link

RECALLED PRODUCT:

Affected Product	NDC #	Lot Number
Accuretic (quinapril HCl/hydrochlorothiazide) 10-12.5 mg Tablet, 90 count	00071-3112-23	FG5379 exp. 08/2024
Accuretic (quinapril HCl/hydrochlorothiazide) 10-12.5 mg Tablet, 90 count	00071-0222-23	EA6686 exp. 04/2022
Accuretic (quinapril HCl/hydrochlorothiazide) 20/12.5 mg Tablet, 90 count	00071-5212-23	FG5381 exp. 08/2024
Accuretic (quinapril HCl/hydrochlorothiazide) 20/12.5 mg Tablet, 90 count	00071-0220-23	EA6665 exp. 04/2022
		CN0640 exp. 04/2022
Accuretic (quinapril HCl/hydrochlorothiazide) 20/25 mg Tablet, 90 count	00071-0223-23	ET6974 exp. 02/2023
Quinapril and Hydrochlorothiazide 20/25 mg Tablet, 90 count	59762-5225-09	FE3714 exp. 02/2023
Quinapril HCl/Hydrochlorothiazide 20/12.5 mg Tablet, 90 count	59762-0220-01	DN6931 exp. 03/2023
		ED3904 exp. 03/2023
		ED3905 exp. 03/2023

Quinapril HCl/Hydrochlorothiazide 20/25 mg Tablet, 90 count	59762-0223-01	DP3414 exp. 02/2023