

Drug Recall

DATE OF RECALL: April 25, 2022

DRUG NAME: Accupril (Quinapril HCl) tablets 10mg, 20mg, 40 mg

RECALLING FIRM: Pfizer

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

FDA LINK: <https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/pfizer-voluntary-nationwide-recall-lots-accuprilr-quinapril-hcl-due-n-nitroso-quinapril-content>

OTHER DETAILS: See FDA Link

RECALLED PRODUCT:

Product Description	NDC Number	Affected Lots	Expiration Date
Accupril (Quinapril HCl) tablets 10 mg, 90 count	00071-0530-23	DR9639	3/31/2023
Accupril (Quinapril HCl) tablets 20 mg, 90 count	00071-0532-23	DX8682	03/31/2023
		DG1188	05/31/2022
Accupril (Quinapril HCl) tablets 40 mg, 90 count	00071-0535-23	DX6031	03/31/2023
		CK6260	05/31/2022