

Drug Recall

DATE OF RECALL: December 22, 2022

DRUG NAME: Quinapril Tablets

RECALLING FIRM: Lupin Pharmaceuticals, Inc.

REASON FOR RECALL: This recall was due to the presence of a nitrosamine impurity, 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/lupin-pharmaceuticals-inc-issues-voluntary-nationwide-recall-four-lots-quinapril-tablets-due>

OTHER DETAILS: N/A

RECALLED PRODUCT:

Product Description	NDC Number	Affected Lots	Expiration Date
Quinapril 20 mg Tablet, 90 count	68180-0558-09	G102929	04/2023
Quinapril 40 mg Tablet, 90 count	68180-0554-09	G100533 G100534 G203071	12/2022 12/2022 03/2024