Quinapril tablets

On December 21, 2022, Lupin Pharmaceuticals Inc. is voluntarily recalling four (4) lots of Quinapril Tablets due to the presence of a nitrosamine impurity, N-Nitroso-Quinapril, observed in recent testing above the Acceptable Daily Intake (ADI) level. To date, Lupin has received no reports of illness that appear to relate to this issue.

Nitrosamines are common in water and foods, including cured and grilled meats, dairy products and vegetables. Everyone is exposed to some level of nitrosamines. These impurities may increase the risk of cancer if people are exposed to them above acceptable levels over long periods of time.

Quinapril tablet USP is an angiotensin-converting enzyme (ACE) inhibitor indicated for the treatment of hypertension, to lower blood pressure. Lowering blood pressure reduces the risk of fatal and nonfatal cardiovascular events, primarily strokes and myocardial infarctions. Quinapril Tablets USP 20mg, and 40mg is packaged in 90 count bottles and was distributed nationwide in the US to wholesalers, drug chains, mail order pharmacies and supermarkets. The recalled lots are included in the table below:

Product	Lot No	Expiry	NDC	UPC	Distribution Dates
Quinapril Tablets USP, 20mg	G102929	04/2023	68180-558-09 (90's)	368180558095	03/15/2021 – 09/01/2022
Quinapril Tablets USP, 40mg	G100533 G100534 G203071	12/2022 12/2022 03/2024	68180-554-09 (90's)	368180554097	

Patients taking, Quinapril Tablets USP, 20mg, and 40mg are advised to continue taking their medication and contact their pharmacist, physician, or medical provider for advice regarding an alternative treatment.

Consumers, wholesalers, distributors, and retailers with questions regarding this recall should contact Inmar Rx Solutions, Inc. at (877) 538-8445 Monday – Friday 09:00 am to 05:00 pm EST. For reimbursement, please have the recalled lots returned to Inmar Rx Solutions, Inc.; the lot number can be found on the side of the bottle label.