Quinapril Tablets – Lupin Pharmaceuticals, Inc.

The FDA announced a consumer level recall of four lots of Lupin's Quinapril tablets due to the presence of a nitrosamine impurity, N-Nitroso-Quinapril, above the acceptable daily intake level.

Product	Quinapril Tablets 20mg Quinapril Tablets 40mg
Recall identification date	12/22/2022
Lot numbers – 20mg	G102929
Lot numbers – 40mg	G100533 G100534 G203071
Manufacturer	Lupin Pharmaceuticals, Inc.
Affected NDCs	68180-558-09, 68180-554-09

Consumers should contact their physician or healthcare provider if they have experienced any problems that may be related to taking or using this product.

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.

Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting program either online,by regular mail or by fax.

- Complete and submit the report **Online:** www.fda.gov/medwatch/report.htm
- **Regular Mail or Fax:** Download form www.fda.gov/MedWatch/getforms.htm or call 1-800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form, or submit by fax to 1-800-FDA-0178
- https://www.fda.gov/safety/report-problem-fda