December 22, 2022

BSE Limited
Department of Corporate Services,
P. J. Towers,
Dalal Street,
MUMBAI - 400 001.

National Stock Exchange of India Limited
Exchange Plaza,
Bandra Kurla Complex,
Bandra (East),
Mumbai - 400 051.

Dear Sir/Madam,

Sub: Disclosure pursuant to Regulation 30 of SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 ('Listing Regulations').

Enclosed is a Press Release as regards voluntary nationwide recall of Quinapril Tablets.

This may kindly be considered as a disclosure pursuant to Regulation 30 of the Listing Regulations.

The above is for your information and dissemination.

Thanking you,

For LUPIN LIMITED

Rajvardhan Vishnu Satam

Digitally signed by Rajvardhan Vishnu Satam
DN: c=IN, st=Maharashtra, 2.5.4.20=c19d8ef961d9fa59e973a093a4d3924b2532eb2f496ea7e2abcf74700f5375b7, postalCode=400104, street=2404,SEJAL TOWER,SEJAL PARK,LINK ROAD,GOREGAON WEST,Mumbai,Motilal Nagar,Borivali,Mumbai Suburban, pseudonym=9a4b7cd219ea516ad56e6d651dcbbe5b, title=0191, serialNumber=df8ebe60a88c796663c738ea25bfa0b475d195f726cdb67a2ff09c4242771b2a, o=Personal, cn=Rajvardhan Vishnu Satam
Date: 2022.12.22 16:00:07 +05'30'

R. V. SATAM
COMPANY SECRETARY
(ACS - 11973)

Encl: a/a

Lupin Pharmaceuticals, Inc. Issues Voluntary Nationwide Recall of Four Lots of Quinapril Tablets due to potential presence of N-Nitroso-Quinapril Impurity

Company Contact:
Shweta Munjal
Email: shwetamunjal@lupin.com

FOR IMMEDIATE RELEASE

Baltimore, Maryland, December 22, 2022: Lupin Pharmaceuticals Inc. is voluntarily recalling four (4) lots of Quinapril Tablets to the patient (consumer/user) level 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.

Lupin discontinued the marketing of Quinapril tablets in September 2022.

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	

Lupin Pharmaceuticals Inc. is notifying its wholesalers, distributors, drug chains, mail order pharmacies and supermarkets by phone and through recall notification and is arranging for the return of all the recalled product lots.

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.

Wholesalers, distributors and retailers that have Quinapril Tablets USP, 20mg, and 40mg that are being recalled should discontinue distribution of the recalled product lots immediately.

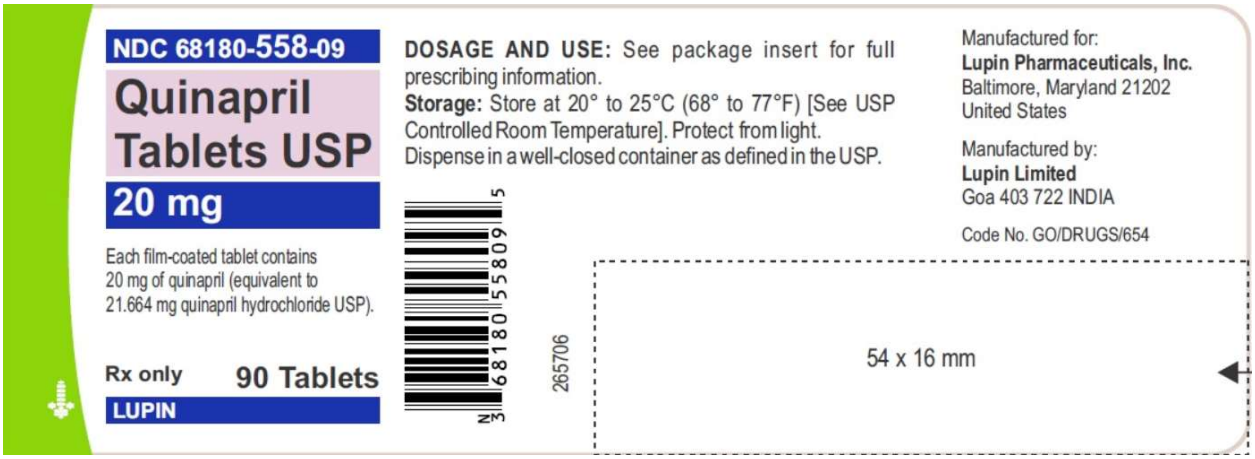
Consumers, wholesalers, distributors, and retailers with questions regarding this recall should contact Inmar Rx Solutions, Inc. at (877) 538-8445 Monday – Friday 09:00 am to 05:00 pm EST. For reimbursement, please have the recalled lots returned to Inmar Rx Solutions, Inc.; the lot number can be found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

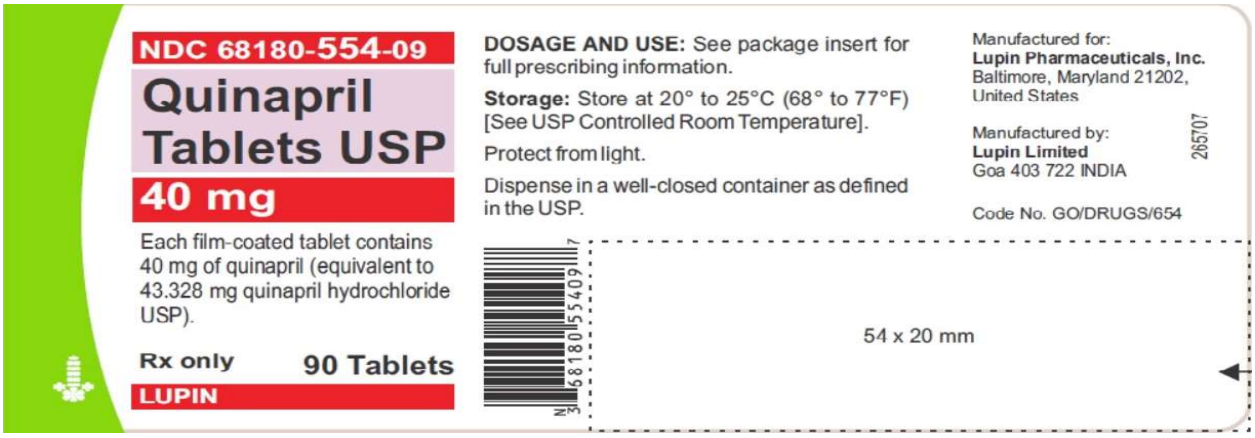
Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting program either online, by regular mail or by fax.

- 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.

This recall is being conducted with the knowledge of the U.S. Food and Drug Administration.

Product Label:





About Lupin Pharmaceuticals

Lupin Pharmaceuticals, Inc. is the U.S. based wholly-owned subsidiary of Lupin Limited and is the 3rd largest pharmaceutical company in the U.S. based on total prescriptions. Together, all Lupin-owned entities combine to make up the 8th largest generic pharmaceutical company in the world by revenue size. Lupin Pharmaceuticals, Inc. is dedicated to delivering high-quality medications across many treatment areas. Lupin Pharmaceuticals Inc.'s branded pharmaceuticals division, is the provider of products designed to help prevent and manage women's health conditions with serious health consequences.