

The Board of Pharmacy has received notice of the following product recall. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

Natco Pharma Limited, Pharma Division, Kothur, Ranga Reddy District, Telangana State, India is initiating a recall to the **RETAIL level** for Product Lansoprazole Delayed-Release Capsules USP, 15 mg (ANDA# A203306) batch # 411988 manufactured by Natco Pharma Limited, India and marketed by Rising Pharma Holdings, Inc. USA.

This recall is based on a complaint received by Natco Pharma Limited from McKesson Global Procurement and Sourcing for batch number 411987. Recall has been initiated for the same complaint batch number. As part of impact assessment, defective capsules were noticed for campaign lot# 411988, therefore, recall has been initiated for the same. Batch# 411988 was shipped to the customers on 01/02/2024.

Natco Pharma Limited, Pharma Division, Kothur, Ranga Reddy District, Telangana, India has conducted a thorough investigation and health hazard assessment. This assessment concluded that Lansoprazole Delayed Release capsules are primarily used to treat hyperacidity of the gastrointestinal tract, inadvertent usage of the defective capsules may potentially lead to subtherapeutic levels of the drug, but no long-term effects or serious hazard to the patient is anticipated.

Product Name	NDC(s)	Lot(s) / Exp. date	Distribution dates
	16571-742-41 (Label)		
	16571-742-42		
Lansoprazole Delayed-Release Capsules USP, 15 mg	(Carton/Shipper label)	411988/MAY 2025	01/02/2024