

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.

Dr. Reddy's Laboratories, Inc. is initiating a recall as of Zoledronic Acid Injection, 5 mg/ 100 ml, Single Dose Vial a precaution based on market complaint, the customer discovered the metal ring around the stopper loosened when the seal was removed causing the vial to leak.

Detectability of this defect before the administration is high and the probability of using such an affected vial by a healthcare professional is very low. However, if an affected vial is used, there could be a suboptimal/ decreased therapeutic response due to a lower dose being administered. Moreover, there is a possibility of contamination of sterile product, which may result in nosocomial infections in the patient, in case the defective vial is actually used.

The product Distribution Dates: March 04, 2024 - April 15, 2024

PRODUCT DESCRIPTION: Zoledronic Acid Injection, 5 mg/ 100 ml, Single Dose Vial

NDC NUMBER: 55111-688-52

LOT NUMBER: G3000010

EXPIRATION DATE: 11/2025