REPUBLIC OF KENYA
MINISTRY OF HEALTH

PHARMACY AND POISONS BOARD

PUBLIC ALERT

VOLUNTARY RECALL OF S-PRAZO (ESOMEPRAZOLE MAGNESIUM DELAYED-RELEASE CAPSULES 40MG) BATCH NO SPZ-302 MANUFACTURED BY LABORATE PHARMACEUTICAL INDIA LIMITED

The Pharmacy and Poisons Board ("the Board") wishes to inform the public of the voluntary recall of S-PRAZO (Esomeprazole Magnesium Delayed-Release Capsules 40mg) **Batch No SPZ-302** Manufactured by Laborate Pharmaceutical India Limited.

The product batch is being recalled following a market complaint regarding mix-up of blister strips; some boxes were found to contain two different blister strips, i.e S-Prazo Capsules and Donystatin Tablets, within the same outer carton.

In view of the above, the Board advises all pharmaceutical outlets, healthcare facilities, healthcare professionals and members of the public to **STOP** further distribution, sale, issuing or use of the product batch and return the specified product batch to their nearest healthcare facility or respective suppliers.

The Board remains committed to protecting the health of the public and encourages the public to promptly report any suspected cases of sub-standard medicines or adverse drug reactions. The reports may be made to the nearest healthcare facility or the Pharmacy and Poisons Board through the following channels:

- Website: https://pv.pharmacyboardkenya.org/users/mpublic
- USSD code at *271#
- Mobile application: mPvERS both Android & iOS
- Email: pms@ppb.go.ke or pv@ppb.go.ke
- Telephone No. 0795 743 049

Dr. F. M. Siyoi
CHIEF EXECUTIVE OFFICER

20th August 2024

PRODUCT PHOTO


S-Prazo
Esomeprazole Magnesium Delayed-Release Capsules USP 40 mg
3x10 Capsules