REPUBLIC OF KENYA
MINISTRY OF HEALTH

PHARMACY AND POISONS BOARD

PUBLIC ALERT

MANDATORY RECALL OF FLAMODIP (AMLODIPINE) 5 MG TABLETS BATCH NO FLD303 MANUFACTURED BY MEDICO REMEDIES PVT LTD

The Pharmacy and Poisons Board ("the Board") has issued a recall for Flamodip (Amlodipine) 5mg Tablets, Batch No. FLD303, manufactured by Medico Remedies Pvt Ltd.

The recall has been initiated due to a labeling error, where the product's label does not accurately reflect its contents. *The secondary packaging is labeled as Flamodip-5 (Amlodipine), while the primary packaging is labeled as Flaminopril-5 (Enalapril).*

In light 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 product to their nearest healthcare facility or respective suppliers.

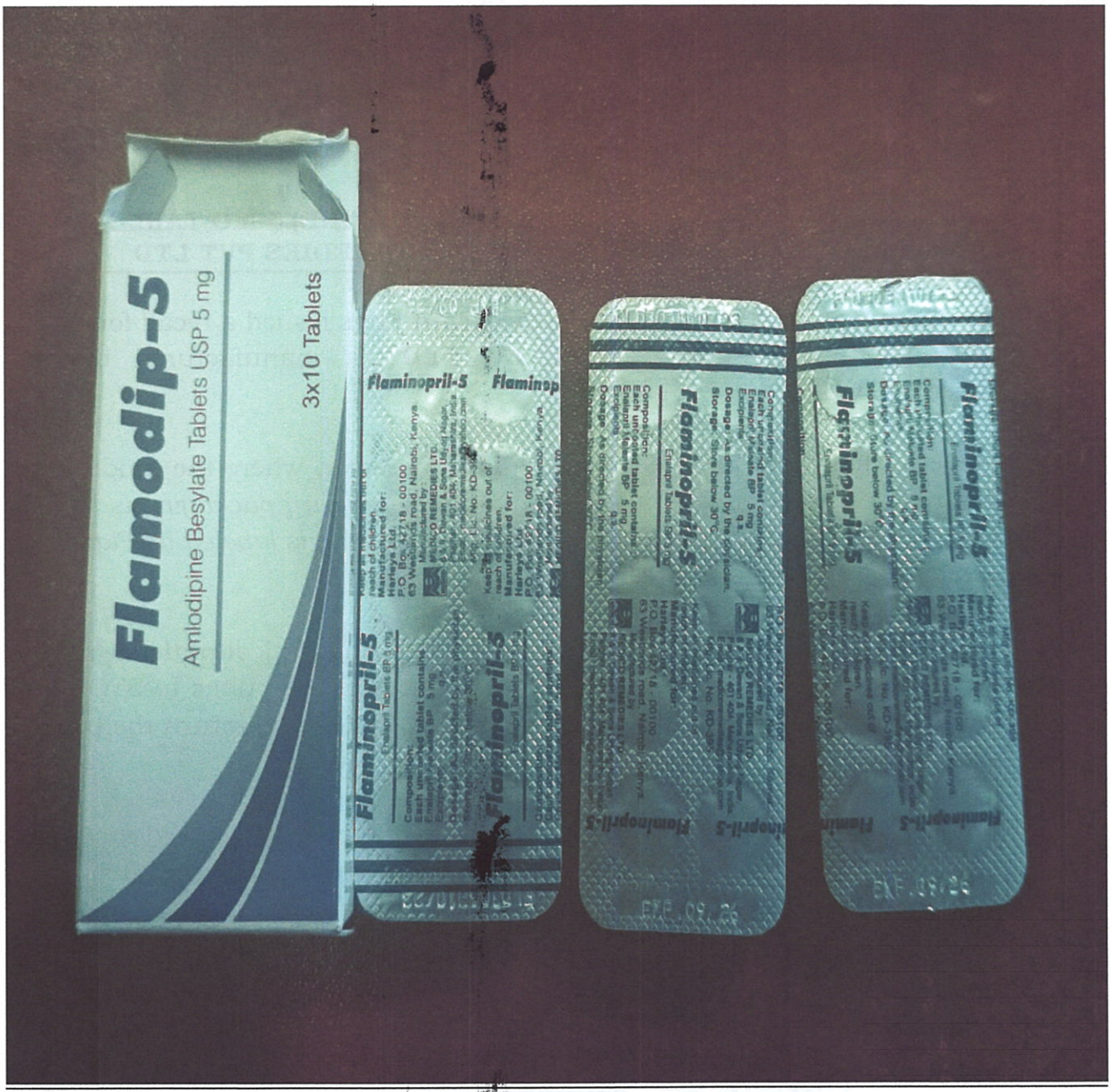
The Board encourages the public to promptly report any suspected cases of sub-standard medicines or adverse drug reactions to the nearest healthcare facility or the Pharmacy and Poisons Board through the following channels:

- https://pv.pharmacyboardkenya.org/users/mpublic
- USSD code at *271#
- Mobile application: mPvERS both Android & iOS
- Email pv@ppb.go.ke or pms@ppb.go.ke
- Telephone No. 0795743049

Dr. F. M. Siyoi
CHIEF EXECUTIVE OFFICER

26th September 2024

PHOTO


Flamodip-5
Amlodipine Besylate Tablets USP 5 mg
3x10 Tablets
Flaminopril-5
Flaminopril-5
Flaminopril-5
Flaminopril-5