TMDA
Tanzania Medicines & Medical Devices Authority

Weekly Surveillance Report





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WSR

"Updates on Substandard, Falsified (SF) and unregistered medical products"

Substandard and falsified medical products can have severe health consequences, including treatment failure, disease progression and prolonged illness, toxicity and harmful adverse effects, increased antimicrobial resistance, loss of public trust in healthcare systems and increased healthcare expenditure and economic losses.

TMDA emphasizes that ensuring the availability of safe, quality and effective medical products is essential for protecting public health in Tanzania.

In lieu to the above emphasis, TMDA implements various strategies to prevent, detect and respond to substandard and falsified medical products through:

- Strengthening regulatory frameworks and enforcement systems;
- Conducting inspections of manufacturers, importers, distributors and healthcare facilities;
- Carrying out post-marketing surveillance and laboratory testing of medical products;
- Monitoring medicine supply chains and distribution systems;
- Issuing safety alerts, recalls and regulatory actions where necessary;
- Collaborating with local and international stakeholders and regulatory authorities;
- Enhancing public awareness on the dangers of purchasing medicines from unauthorized sources; and
- Promoting the use of technological solutions and traceability systems to improve monitoring and detection.
- During the period lasting from 3rd to 9th August, 2026, inspectors from the Tanzania Medicines and Medical Devices Authority (TMDA) at the Southern Highland Zone successfully intercepted eight (8) unregistered medicines during the routine Post Marketing Inspection. Details of the unregistered medicinal products are presented in table 1 below.



Table 1: List of unregistered Medicines Seized at Southern Highland Zone from 3rd to 9th August, 2026

S/N	Name of the Product (Brand and generic)	Batch details	Name of Manufacturer	Quantity and dosage form
1.	Levobact 750 (Levofloxacin Hemihydrate tablets 750)	LFEH0039	Micro labs LIMITED HOSUR INDIA	10 Tablets
2.	VEGA 100 Sildenafil citrate	-	Signature	38 tabs
3.	Contrasafe (medroxyprogesterone acetate)	4003343	Senador Laboratory Pvt Ltd Gujarat INDIA.	03 vials
4.	Heparin Sodium Injection	HPV300924	Questus Pharma private Ltd Telangana INDIA.	1 vial
5.	INDOREN (Indomethacin capsules)	01824	Rene Industries Ltd Kampala Uganda.	190 capsules
6.	Caberlee (Carbegoline tablets)	T242574	Johnlee Pharmaceuticals Pvt. Ltd., India.	32 tablets
7.	FLUZICAN (Fluphenazine decanoate Injction)	GN7886	American Remedies, India.	10 vials
8.	SYNDOPA (Levodopa and carbidopa tablts)	GKF112A	Sun pharma Laboratories Ltd. Gangtok India	420 tabs

Photo of the selected unregistered products reported from Southern Highland Zone are presented below as Figure 1 to 3

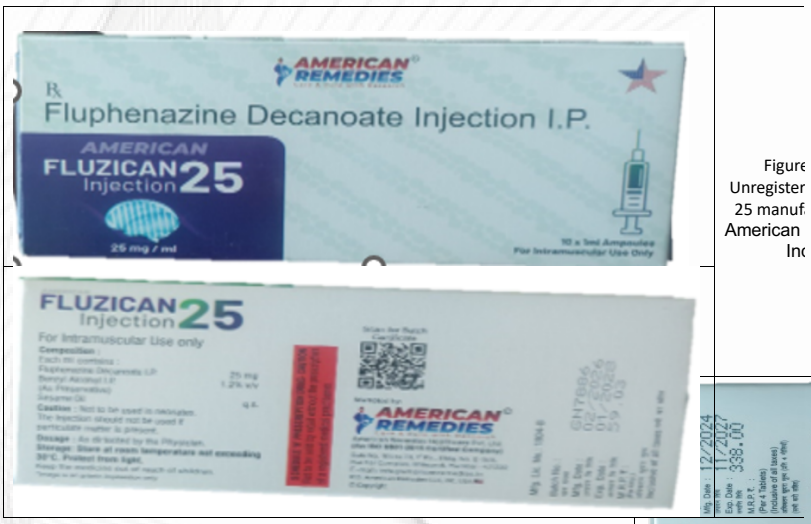
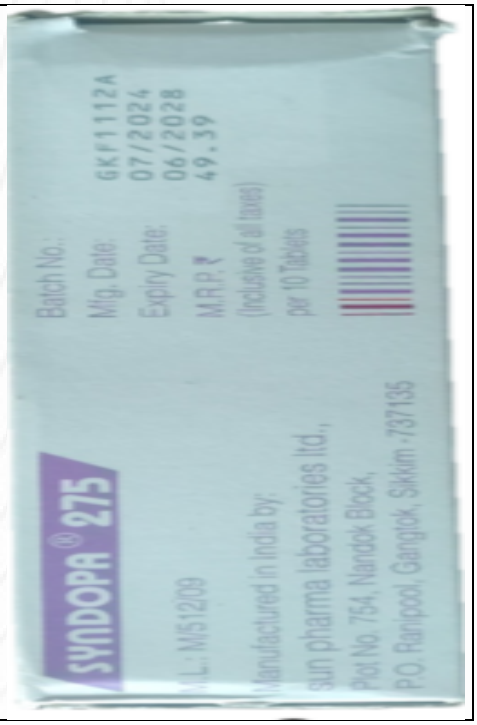


Figure
Unregister
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Figure No. 1: **Unregistered Caberlee-0.5 manufactured by** Johnlee Pharmaceuticals Pvt. Ltd., India

Figure No. 3: Unregistered Syndopa 275 manufactured by Sun pharma Laboratories Ltd. Gangtok India.		

All unregistered medicines were confiscated and will be disposed at owner's cost. Fines were imposed for those who were found with unregistered products.

The Authority continues to warn the public against purchasing or consuming any medicinal product that does not appear in the official register. If you encounter any suspicious or unregistered products, they should be reported immediately to the nearest TMDA office or via the toll-free number **0800110084.**



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