MEDICAL PRODUCT ALERT

DRAP ALERT NO. Nº I/S/12-24-57

RECALL NOTICES ISSUED BY PROVINCIAL DRUG TESTING LABORATORIES.

Date: 31st December, 2024.

Target Audience:

- National Regulatory Field Force of DRAP and Provincial Drug Control Departments.
- Healthcare Professionals (Physicians, Pharmacists & Nurses) & Veterinary professionals.
- General Public.

Alert Summary:

DDCP PV Lahore and Secretary PQCB Quetta has informed DRAP regarding recall of following batches of mentioned products:

S#	Product Name	Batch No.	Manufactured by	Test Results
1	Injection I-Vitamin Super 100ml (**FOR VET USE ONLY**) (Each ml contains: sodium Selenite 0.5mg, Vit. E 70mg, Vit. B12 0.1mg, Vit. B1 20mg, Adenosine-5 Monophosphate 5mg) Reg# 062066	6501	International Pharma Labs, Raiwind Road, Bhobtian Chowk, Defence road, 1Km towards Kahna, Lahore.	Adulterated and Substandard
2	Injection Oxytocin 50ml (**FOR VET USE ONLY**) (Oxytocin 10IU/ml) Reg# 020830	OXY-2219	Lawrance Pharma (Pvt.) Ltd., 10.5 Km Shekhupura road, Lahore.	Substandard
3	Injection Flagyl 100ml (Metronidazole 500mg/100ml) Reg# 005102	AD354	Sanofi-Aventis Pakistan Limited, Plot no. 23, Sector 22, Korangi Industrial Area, Karachi.	Substandard
4	Capsule Esopase 20mg (Esomeprazole 20mg) Reg# 045914	3K159	Lucky Core Industries Ltd., 32/2A, Phase 3, Industrial Estate Hattar	Substandard

Action initiated: -

The regulatory field force of DRAP and Provincial Drug Control departments have been requested to immediately conduct market surveys for detection and removal of these products from the market.

Advice for Healthcare Professionals: -

DRAP requests increased vigilance within the supply chains of institutions/pharmacies/healthcare

facilities likely to be affected by batches of mentioned products. Adverse reactions or quality problems experienced with the use of these products are to be reported to the National Pharmacovigilance Centre (NPC), DRAP using Adverse Event Reporting Form or online through this link. Further information of reporting problems to DRAP is available on this link.

Advice for Consumers / general public: -

Consumers should stop using products bearing the affected batch number(s) and shall contact to their physician or healthcare provider if they have experienced any problems that may be related to taking or using this drug product, and report the incident to Drug Regulatory Authority of Pakistan/ National Pharmacovigilance Centre.

Drug Regulatory Authority of Pakistan
محفوظ، موثر اور معیاری اشیائے علاج

DRAP, Islamabad
+92 051 9255969
gsms@dra.gov.pk