

## DRUG REGULATORY AUTHORITY OF PAKISTAN DIVISION OF QUALITY ASSURANCE & LABORATORY TESTING

### **MEDICAL PRODUCT ALERT**

**DRAP ALERT NO.** Nº I/S/10-25-104

### **VETERINARY DRUG PRODUCTS DECLARED SUBSTANDARD**

BY PROVINCIAL DRUG LABORATORIES, KARACHI.

Date: 30<sup>th</sup> October, 2025

### **Target Audience:**

- National Regulatory Field Force of DRAP and Provincial Drug Control Departments.
- Healthcare Professionals-Veterinarians
- Farmers/consumers

### **Alert Summary:**

Drugs Testing Laboratories from Provinces informed that the sample of below mentioned **Veterinary products** have been declared as *'Substandard / Misbranded'*.

S#	Product Details	Batch #	Manufacturer details	Remarks
1.	Inj. Genta-Combisone 50ml  Each 100 ml contains: Tylosin Tartrate 15 gm Gentamicin Sulphate6 gm Dexamethasone0.0265 gm Chlorpheniramine 0.750 gm (Reg # 046696)	GC-168	M/s Leads Pharma (Pvt) Ltd. Plot No. 81-A Street No. 6 I- 10/3 Islamabad. (DML # 000392)	The sample is <b>Sub-Standard</b> with regards to Sterility Test.
2.	Injection Ivergen 10ml Each ml contains: Ivermectin10 mg (Reg. # 023432)	594.IV	M/s Symans Pharmaceuticals (Pvt) Ltd. 10-Km Sheikhupura Road Lahore. (DML # 000323)	The sample is <b>Sub-Standard</b> with regards to Assay and Test for related substances.

### **Risk Statement:**

The use of these defective **veterinary injectable products** may adversely affect livestock and farm animals, particularly those treated for infections or parasitic infestations. *Injection Genta-Combisone* poses a risk of microbial contamination due to sterility failure, which can lead to abscesses, septicemia, and reduced milk or meat yield in cattle, buffalo, goats, and other farm species. *Injection Ivergen*, being sub-standard in assay and related substances, may result in incomplete parasite control, therapeutic failure, and possible drug resistance in treated herds. These defects ultimately endanger the **health and productivity of livestock**, causing **economic losses to farmers** and potentially impacting the **safety of animal-derived food products** consumed by the general public.









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### **Action initiated: -**

The field force of DRAP and Provincial Drug Control departments have been directed to immediately conduct market surveys for detection of presence and removal of mentioned batches from the market.

### Advice for Veterinarian: -

This alert applies strictly to the specific batch listed above and does not apply to other batches of the **same** product. Veterinarians are therefore advised not to prescribe, administer, or stock these identified batch under any circumstances.

#### **Advice for Farmers and Livestock Owners:**

Farmers and livestock owners are advised **not to use** the affected batches of *Injection Genta-Combisone* (*Batch* # *GC-168*) and *Injection Ivergen* (*Batch* # *594.IV*) on their animals. The use of non-sterile or sub-standard veterinary injections can cause **serious illness**, **swelling**, **infection**, **or treatment failure** in cattle, buffalo, goats, and other livestock.

If any animals have recently been treated with these products and show signs of **fever**, **swelling at the injection site**, **weakness**, **or loss of appetite**, stop using the product immediately and **consult a qualified veterinarian**.

Farmers should **return the remaining stock** to the supplier, distributor, or the nearest veterinary office. They are further advised to purchase veterinary medicines only from **authorized dealers** and ensure that the products bear **clear batch numbers, expiry dates, and registration details** issued by DRAP.

For continued livestock health and to avoid losses, always **store veterinary medicines properly**, follow veterinary guidance for dosage, and report any suspected quality-related issues to the **Provincial Livestock Department or DRAP** through the official complaint channels.







