DRUG REGULATORY AUTHORITY OF PAKISTAN
Division of Quality Assurance and Laboratory Testing

RAPID ALERT

DRAP ALERT No: Nº I/S/09-25-70

SPURIOUS / FALSIFIED MEDICINE DETECTED – DAIZELIVE INJECTION 2ML (DIAZEPAM 10MG/2ML, CLAIMED)

Date: 17th September , 2025

Target Audience:

- Regulatory Field Force of DRAP and Provincial Drug Control departments.
- Healthcare Professionals
- Consumers

Problem Statement:

Directorate of Drugs Control Punjab informed Drug Regulatory Authority of Pakistan that the samples of below mentioned products have been declared as '***Spurious***'.

S#	Product Name	Batch No.	Manufacturers	Remarks
1.	**Injection Daizelive 2ml** Each 2ml ampoule contains: Diazepam …... 10 mg	DZ44	**M/s Liven Pharmaceuticals (Pvt) Ltd.** Sray Road, 49 KM, Multan Road, Phool Nagar, District Kasur (DML # 000881)	Drug Testing Laboratory, Rawalpindi has declared the sample as **'spurious'** since it does not contain the claimed active ingredient (Diazepam) and instead contains an undeclared ingredient (Chlorpheniramine Maleate 45.283 mg/2ml).

Risk Statement:

Laboratory testing has confirmed that the product labeled as **Daizelive Injection 2ml (Diazepam 10mg/2ml)** is **spurious/falsified**, as it does not contain **Diazepam**, instead contains **Chlorpheniramine Maleate (45.283mg/2ml)**.

Use of this falsified product poses a **serious and life-threatening risk**, particularly for patients requiring Diazepam for **seizures, severe anxiety, or emergency care**, as they will not receive the intended treatment. Healthcare professionals and the public are strongly advised to **immediately discontinue use of this product**, ensure recall compliance, and report any suspected adverse outcomes or suspicious stock to DRAP.

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

Advice for Healthcare Professionals: -

DRAP requests healthcare professionals to have increased vigilance within the supply chains of institutions/pharmacies/healthcare facilities likely to be affected by above mentioned product. Any adverse events or quality problems experienced with the use of these products should be reported to National or Provincial pharmacovigilance centers using Adverse Event Reporting Form or online through this link. Further information on reporting problems to DRAP is available on this link.

Advice for Consumers:

Consumers should not use these products and should contact their physician or healthcare provider(s) if they have experienced any problem related to taking or using the above mentioned products and should report the incident to the Drug Regulatory Authority of Pakistan / National Pharmacovigilance Centre. All therapeutic goods must be obtained from licensed pharmacies and other authorized/licensed retail outlets. The authenticity and condition of products should be carefully checked. Seek advice from your pharmacists or other healthcare professionals in case of any doubt.

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

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