Government of Pakistan
Ministry of National Health Services
Regulations & Coordination
Drug Regulatory Authority of Pakistan

Sr. No. 4394

(For Government Supply/Institutions only)

Certificate No. F. 3-13/2018-Addl. Dir. (QA & LT-I)-14 **Date of Issue:** 15th February, 2022

CERTIFICATE OF GOOD MANUFACTURING PRACTICES

It is certified that **M/s Davis Pharmaceutical Laboratories, Plot No. 121, Industrial Triangle, Kahuta Road, Islamabad-Pakistan** holding Drug Manufacturing License Number **000432** is authorized to produce drugs/medicine (s). I certify that the site indicated on this certificate complies with current Good Manufacturing Practices (cGMP) in terms of process control, maintenance of equipments, documentation and areas etc, as per provision of Drugs Act, 1976 and rules framed there under. The dosage forms and activities in the following categories:-

Dosage form(s)	Pharmaceuticals Category(ies)	Activity(ies)
Tablet	• *General*	*Mixing, Granulation, Drying, Compression, Coating and Packaging*
Capsule	• *General*	*Mixing, Drying, Filling & Packing*
Capsule	• *Cephalosporin*	*Mixing, Drying, Filling & Packing*
Liquid Syrup	• *General*	*Mixing, Filling & Packing*
Cream/ Ointment/ Gel	• *General*	*Mixing, Filling & Packing*
Sachet	• *General*	*Mixing, Drying, Filling & Packing*
Dry Suspension	• *Cephalosporin*	*Mixing, Drying, Filling & Packing*

2. This certificate is based on the inspection and evaluation conducted on **02-02-2022**

- This certificate is valid till **01-02-2024** or till revoked by this Authority.
- The responsibility to maintain quality as per standards of Good Manufacturing Practices throughout the period of validity of this certificate in manufacturing process of the individual batches of the pharmaceuticals products lies with the manufacturer.
- This certificate permits the firm to apply for registration of their products, manufactured as per valid current good manufacturing practices (cGMP) under Drugs Regulatory Authority of Pakistan, in Pakistan.
- The validity will automatically seize in case of reporting of non-compliance of current Good Manufacturing Practices (cGMP) under the Drugs Act, 1976 and rules framed there under.
- This certificate is in line with the format as recommended by WHO (TRS No. 908, 2003).
- This certificate is issued on the demand of M/s Davis Pharmaceuticals Lab, Islamabad, for Government supply/institutions only.

Name of Certifying Authority: **Dr. Hafsa Karam Elahi**
Designation: **Additional Director (QA & LT-I), DRAP, Islamabad**
Address of certifying Authority: 3rd Floor, Telecom Foundation Complex, Mauve Area, G-9/4, Kashmir Highway, Islamabad.

Signature ----------

Digitally signed by Chitic Vlad
Date: 2022.06.02 21:00:22 EEST
Reason: MoldSign Signature
Location: Moldova

Stamp

Drug Regulatory Authority of Pakistan
Additional Director
Division of QA&LT
Islamabad
Government of Pakistan