Government of Pakistan
Ministry of National Health Services, Regulations & Coordination
Drug Regulatory Authority of Pakistan
2nd Floor, USAID Building, No IV, Block B, SMCHS, Karachi
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CERTIFICATE OF GOOD MANUFACTURING PRACTICES

Certificate No: **GMP-C-0CAB9D** Date of Issue: **14/10/2025**

It is certified that **AGP LIMITED B-23-C, SINDH INDUSTRIAL TRADING ESTATE, (SITE), Karachi, Karachi West Site Town** holding Drug Manufacturing License Number **000348** is authorized to product 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:-

Sr #	Dosage Form(s)	Pharmaceuticals Category(ies)	Activities
1	Tablet	General	Mixing, Drying, Granulation, Compression, Coating and Packaging.
2	Capsule	General	Mixing, Drying, Filling and Packaging
3	Sachet	General	Mixing, Sachet Filling, Sealing and Packaging
4	Oral Liquid	General	Mixing, Filling and Packaging
5	Semi solid (Cream/Ointment)	General	Mixing, Filling and Packaging
6	Liquid Ampoule	General	Mixing, Filling and Packaging
7	Dry Powder Suspension	General	Mixing, Filling and Packaging

This certificate is based on the inspection and evaluation conducted on **04/08/2025**

This certificate is valid for 3 years till **14/10/2028** 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 manufacture.

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 ease 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, 2023).

This certificate is issued on the demand of AGP LIMITED

The Certificate is valid only within Pakistan.

Officer Remarks:
Approved

Name of Certifying Authority: **Abdul Rasool Shaikh**

Address of Certifying Authority: 2nd Floor, USAID Building, No IV, Block B, SMCHS, Karachi

Note: This is the system generated document and does not require any signature. This document can be verified electronically/Online the QR code available on the document.