

	<b>ACCREDITATION DOCUMENT</b>	<b>F-06/02</b> <b>Issue Date: 18/08/2020</b> <b>Rev. No: 09</b> <b>LAB 107</b>
---	-----------------------------------	---

**Accreditation No: LAB 107**

**Awarded to**

**Bio-Analytical / Pharmaceutical QC Laboratory  
Institute of Biological, Biochemical and Pharmaceutical Sciences.  
Karachi-Pakistan.**

The scope of accreditation is in accordance with the standard specifications outlined in the following page(s) of this document. The accredited scope shall be visible and legible in areas such as customer service, sample-receiving section etc and shall not mislead its users.

The accreditation was first time granted on **21-07-2016** by Pakistan National Accreditation Council.

The laboratory complies with the requirements of **ISO/IEC 17025:2017**.

The accreditation requires regular surveillance, and is valid until **20-07-2028**.

The decision of accreditation made by Pakistan National Accreditation Council implies that the organization has been found to fulfill the requirements for accreditation within the scope.

The organization however, itself is responsible for the results of performed measurements/tests.

**PAKISTAN NATIONAL ACCREDITATION COUNCIL**

28-07-2025

Date

SD

Director General

	<b>ACCREDITATION DOCUMENT</b>	<b>F-06/02</b> <b>Issue Date: 18/08/2020</b> <b>Rev. No: 09</b> <b>LAB 107</b>
---	-----------------------------------	---

### **Testing Laboratory.**

**Accreditation Scope of IBBPS Bio Analytical / Pharmaceutical QC Laboratory,**  
**TLA Old Building 1st Floor DUHS Ojha Campus, Karachi, Pakistan.**

#### **Permanent laboratory premises**

<b>Materials/Products tested</b>	<b>Testing field (e.g. environmental testing or mechanical testing)</b>	<b>Types of test/ Properties measured /technique</b>	<b>Reference to standardized method (e.g. ISO 14577-1:2003)/ Internal method reference</b>
Clarithromycin Suspension	Drug Testing	Drug Assay	HPLC (USP Method: PTL/STP/CLA/007)
Cefadroxil 500mg Capsule	Drug Testing	Drug Assay	HPLC (USP Method: PTL/STP/CEF/026)
DEG (Diethylene Glycol) & EG (Ethylene Glycol)	Drug Testing	Impurity Testing	GC-FID (USP Method: PTL/STP/EGDG/004 for raw materials)&In-House Method:(PTL/STP/EGDEG/005 for finished dosage forms
NDMA & NDEA	Drug Testing	Impurity Testing	GCMS(In House Method: PTL/STP/NDMANDEA/002

28-07-2025

Date

Sd

Director