Accreditation No: LAB 215

Awarded to

Magns Pharmaceuticals QC Laboratory
Plot # 7B Value Addition City Sahianwala Road Khurrianwala Faisalabad, 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 **20-01-2021** 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 **19-01-2024.**

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

20-01-2021
Date

Sd
Director General

Testing Laboratory.

Accreditation Scope of Magns Pharmaceuticals QC Laboratory
Plot # 7B Value Addition City Sahianwala Road Khurrianwala
Faisalabad, Pakistan

Permanent laboratory premises X

Materials/Products tested	Testing field (e.g. environmental testing or mechanical testing)	Types of test/ Properties measured	Reference to standardized method (e.g. ISO 14577-1:2003)/ Internal method reference
Cflox Tablet (Ciprofloxacin as HCl)	Pharmaceutical Testing (Tablet)	Assay	USP 2020 Vol. I Page 1015 Chromatography HPLC
		Dissolution Test	USP 2020 Vol. I Page 1015 UV spectrophotometer
		Disintegration Test (Film Coated)	USP 2020 Vol. VII Page 6940 (701) DT Apparatus
		Friability (Uncoated Tablet)	USP 2020 Vol. VIII Page 8137 (1216) Friability Apparatus
		Average Weight /Weight Variation,	USP 2020 Vol. VII Page 7183 (905) Weighing Balance
		Average Thickness (Uncoated Tablet)	Internal Method QC/AP/FG/005 Vernier Calliper
Leufex Tablet (Levofloxacin as Hemihydrate)	Pharmaceutical Testing (Tablet)	Assay	USP 2020 Vol. III Page 2609 Chromatography HPLC
		Dissolution Test	USP 2020 Vol. III Page 2609 UV spectrophotometer
		Disintegration Test (Film Coated)	USP 2020 Vol. VII Page 6940 (701) DT Apparatus
		Friability (Uncoated Tablet)	USP 2020 Vol. VIII Page 8137 (1216) Friability Apparatus
		Average Weight /Weight Variation,	USP 2020 Vol. VII Page 7183 (905) Weighing Balance
		Average Thickness (Uncoated Tablet)	Internal Method QC/AP/FG/006 Vernier Calliper

20-01-2020
Date

Sd
Director

Materials/Products tested	Testing field (e.g. environmental testing or mechanical testing)	Types of test/ Properties measured	Reference to standardized method (e.g. ISO 14577-1:2003)/ Internal method reference
Veloft (Delayed Release Tablet) Diclofenac Sodium	Pharmaceutical Testing (Tablet)	Assay	USP 2020 Vol. II Page 1348 Chromatography HPLC
		Dissolution Test	USP 2020 Vol. II Page 1349 UV spectrophotometer
		~~Disintegration Test~~	USP 2020 Vol. VII Page 6940 (701) DT Apparatus
		Friability (Uncoated Tablet)	USP 2020 Vol. VIII Page 8137 (1216) Friability Apparatus
		Weight Variation,	USP 2020 Vol. VII Page 7183 (905) Weighing Balance
		Average Thickness & Average Diameter (coated)	Internal Method QC/AP/FG/029 Vernier Calliper
Czet Tablet (Cetirizine 2HCl)	Pharmaceutical Testing (Tablet)	Assay	BP 2020 Vol. III Page 324 Chromatography HPLC
		Dissolution Test	BP 2020 Vol. III Page 324 UV spectrophotometer
		Disintegration Test (Film Coated)	BP 2020 Vol. V A384 DT Apparatus
		Friability (Uncoated)	BP 2020 Vol. V A572 Friability Apparatus
		Average Weight /Weight Variation,	BP 2020 Vol. V A408 Weighing Balance
		Average Thickness & Average Diameter (Coated)	Internal Method QC/AP/FG/021 Vernier Calliper
Veloft (Prolong Release Capsule) Diclofenac Sodium	Pharmaceutical Testing (Capsule)	Assay	BP 2020 Vol. III Page 506 UV spectrophotometer
		Average Weight /Weight Variation,	BP 2020 Vol. V A408 Weighing Balance

20-01-2020
Date

Sd
Director

Materials/Products tested	Testing field (e.g. environmental testing or mechanical testing)	Types of test/ Properties measured	Reference to standardized method (e.g. ISO 14577-1:2003)/ Internal method reference
Veloft (Extended Release Tablet) Diclofenac Sodium	Pharmaceutical Testing (Tablet)	Assay	USP 2020 Vol. II Page 1350 Chromatography HPLC
		Dissolution Test	USP 2020 Vol. II Page 1350 Table 3 UV spectrophotometer
		Friability (Uncoated)	USP 2020 Vol. VIII Page 8137 (1216) Friability Apparatus
		Weight Variation,	USP 2020 Vol. VII Page 7183 (905) Weighing Balance
		Average Thickness & Average Diameter (Coated)	Internal Method QC/AP/FG/030 Vernier Calliper

20-01-2020
Date

Sd
Director