



ACCREDITATION DOCUMENT

F-06/02
Issue Date: 10/08/15
Rev. No: 07
LAB 062

Accreditation No: LAB 062

Awarded to

**Quality Control Laboratory
Saffron Pharmaceuticals (Pvt.) Ltd.,
19K-m Sheikhupura Road, 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 **08-04-2013** 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 **22-09-2020**.

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

06-01-2020

Date

Director General



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Testing Laboratory.

Accreditation Scope of Quality Control Laboratory, Saffron Pharmaceuticals (Pvt.) Ltd., Faisalabad, Pakistan.

Permanent laboratory premises

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
Lucid 250mg Tablet	Pharmaceutical Chemical testing	Assay	BP(2020)/HPLC, Alternate Internal Method/ UV-Vis. Spectrophotometer
	Pharmaceutical Physical testing	Disintegration	BP(2020) Appendix XII A. Disintegration Test for Tablets and Capsules / Disintegration Apparatus
	Pharmaceutical Physical testing	Identification Test	BP(2020)/HPLC, Alternate Internal Method/ UV absorbance / UV-Vis. Spectrophotometer
	Pharmaceutical Physical testing	Uniformity of weight	BP(2020) Appendix XII G. Uniformity of Weight (Mass) / Analytical Balance
Lucid 500mg Tablet	Pharmaceutical Chemical testing	Assay	BP(2020)/HPLC, Alternate Internal Method/ UV-Vis. Spectrophotometer
	Pharmaceutical Physical testing	Disintegration	BP(2020) Appendix XII A. Disintegration Test for Tablets and Capsules / Disintegration Apparatus
	Pharmaceutical Physical testing	Identification Test	BP(2020)/HPLC, Alternate Internal Method / UV absorbance / UV-Vis. Spectrophotometer

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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
	Pharmaceutical Physical testing	Uniformity of weight	BP(2020) Appendix XII G. Uniformity of Weight (Mass) / Analytical Balance
Locus 250mg Tablet	Pharmaceutical Chemical testing	Assay	USP 42 (NF 37) /HPLC, Alternate Internal Method/UV Vis. Spectrophotometer
	Pharmaceutical Physical testing	Disintegration	USP 42 (NF 37) <701> Disintegration /DT Apparatus
	Pharmaceutical Physical testing	Identification Test	USP 42 (NF 37) /HPLC Alternate Internal Method / UV absorbance / UV Vis. Spectrophotometer
	Pharmaceutical Physical testing	Uniformity of weight	USP 42 (NF 37) <905> Uniformity of dosage units/Analytical Balance
Locus 500mg Tablet	Pharmaceutical Chemical testing	Assay	USP 42 (NF 37) /HPLC, Alternate Internal Method/UV Vis. Spectrophotometer
	Pharmaceutical Physical testing	Disintegration	USP 42 (NF 37) <701> Disintegration /DT Apparatus
	Pharmaceutical Physical testing	Identification Test	USP 42 (NF 37) /HPLC Alternate Internal Method/ UV absorbance / UV Vis. Spectrophotometer
	Pharmaceutical Physical testing	Uniformity of weight	USP 42 (NF 37) <905> Uniformity of dosage units / Analytical Balance
Ofloban 200mg Tablet	Pharmaceutical Chemical testing	Assay	USP 42 (NF 37) /HPLC, Internal / UV Vis. Spectrophotometer

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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
Locus infusion 500mg/100mL	Pharmaceutical Physical testing	Disintegration	USP 42 (NF 37) <701> Disintegration /DT Apparatus
	Pharmaceutical Physical testing	Identification Test	USP 42 (NF 37) / Liquid chromatography /HPLC, Internal / UV absorbance / UV Vis. Spectrophotometer
	Pharmaceutical Physical testing	Uniformity of weight	USP 42 (NF 37) <905> Uniformity of dosage units / Analytical Balance
Mitonil Cream 5 % w/w	Pharmaceutical Chemical testing	Assay	Internal / UV absorbance / UV Vis. Spectrophotometer
	Pharmaceutical Physical testing	pH	BP (2020) Appendix V L. Determination of pH Values / pH meter
	Pharmaceutical Physical testing	Identification Test	Internal / UV absorbance / UV Vis. Spectrophotometer
	Pharmaceutical Microbiological testing	Sterility Test	USP 42 (NF 37) <71> Sterility Test/ LFC, incubators, Filtration assembly
	Pharmaceutical Microbiological testing	Bacterial Endotoxin Test	USP 42 (NF 37) <85> Bacterial Endotoxin Testing / Chromogenic method / UV Vis. Spectrophotometer
Mitonil Cream 5 % w/w	Pharmaceutical Chemical testing	Assay	Internal / Liquid chromatography /HPLC
	Pharmaceutical Physical testing	Identification Test	Internal / Liquid chromatography /HPLC

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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
	Pharmaceutical Physical testing	pH	BP (2020) Appendix V L. Determination of pH Values / pH meter
Bezel Tablet 500 µg	Pharmaceutical Chemical testing	Assay	Internal / Liquid chromatography /HPLC
	Pharmaceutical Physical testing	Disintegration	BP (2020) Appendix XII A. Disintegration Test for Tablets and Capsules / Disintegration Apparatus
	Pharmaceutical Physical testing	Identification Test	Internal / Liquid chromatography /HPLC
	Pharmaceutical Physical testing	Uniformity of weight	BP (2020) Appendix XII G. Uniformity of Weight (Mass) / Analytical Balance
Bezel Injection 500 µg /mL	Pharmaceutical Chemical testing	Assay	Internal / UV-Vis Spectrophotometer
	Pharmaceutical Physical testing	pH	BP (2020) Appendix V L. Determination of pH Values / pH meter
	Pharmaceutical Physical testing	Identification Test	Internal / UV absorbance / UV-Vis. Spectrophotometer
	Pharmaceutical Microbiological testing	Sterility Test	USP 42 (NF 37) <71> Sterility Test/ LFC, incubators, Filtration assembly
	Pharmaceutical Microbiological testing	Bacterial Endotoxin Test	USP 42 (NF 37) <85> Bacterial Endotoxin Testing / Chromogenic method / UV Vis. Spectrophotometer
Elgin Tablet 500 µg	Pharmaceutical Chemical testing	Assay	Internal / Liquid chromatography / HPLC

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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
	Pharmaceutical Physical testing	Disintegration	BP (2020) Appendix XII A. Disintegration Test for Tablets and Capsules / Disintegration Apparatus
	Pharmaceutical Physical testing	Identification Test	Internal / Liquid chromatography / HPLC
	Pharmaceutical Physical testing	Uniformity of weight	BP (2020) Appendix XII G. Uniformity of Weight (Mass) / Analytical Balance

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