

ACCREDITATION CERTIFICATE

Issued under the authority of Bangladesh Accreditation Act, 2006 by Bangladesh Accreditation Board (BAB), Ministry of Industries to

Quality Control Laboratory

Drug International Limited (Unit-2), 13/A, 14/A, Tongi I/A

Gazipur, Bangladesh

This is to certify that this

Testing Laboratory

is accredited in accordance with the international standard

ISO/IEC 17025:2017

in respect of the associated scope, subject to the terms and conditions governing the relevant conformity assessment body (CAB) accreditation.

Certificate Number

01.066.22

Accreditation Date

01 March 2022

Date of Issuance

01 March 2022

Date of Expiration

28 February 2025





Md. Monwarul Islam Director General

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This certificate must be returned on request; reproduction must follow BAB guidelines. For the specific scopes to which this accreditation applies, please refer to the Directory of CABs at BAB website.



BANGLADESH ACCREDITATION BOARD (BAB)

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ACCREDITATION SCHEME

(For Testing Laboratory)

CAB Name & Address:

Quality Control Laboratory

Drug International Limited (Unit -2), 13/A, 14/A Tongi I/A, Gazipur, Bangladesh

Accreditation Standard:

ISO/IEC 17025:2017

Accreditation Date:

01 March 2022

Certificate Number:

01.066.22

Issued on:

01 March 2022

Last Amended on:

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Valid until:

28 February 2025

Amendment n	O:

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S.N.	Products/ Materials/ Items of test	Type of tests performed	Specifications/ Standard test methods/Techniques used	Range of testing/Limit of detection
Field:	Chemical (Pharm	aceutical)		
01.	Amoxicillin Trihydrate API	Identification by Infrared Absorption Spectrophotometry (IR)	BP 2018, Volume-I	650-4000cm ⁻¹
		Water Determination by Titration (KF)	BP 2018, Volume-I	0.1% to 100%
		Assay by Liquid Chromatography (HPLC)	BP 2018, Volume-I	0.016mg/ml
02.	Flucloxacillin Sodium API	Identification by Infrared Absorption Spectrophotometry (IR)	BP 2018, Volume-I	650-4000cm ⁻¹
		Water Determination by Titration (KF)	BP 2018, Volume-I	0.1% to 100%
		Assay by Liquid Chromatography (HPLC)	BP 2018, Volume-I	0.044mg/ml
03.	Capsule Demoxil 500 (Amoxicillin Trihydrate BP)	Assay by Liquid Chromatography (HPLC)	BP 2018, Volume-III	0.07mg/ml
04.	Capsule Flupen 500 (Flucloxacillin Sodium BP)	Assay by Liquid Chromatography (HPLC)	BP 2018, Volume-III	0.007mg/ml
Field:	Microbiology	**		
01.	Pharmaceutical	Total Aerobic microbial count	USP 42,(61)	≥ 10 CFU/g
	Finish Product and Excipient	Total yeast & mold count	USP 42,(61)	≥ 10 CFU/g
		Detection of Escherichia coli	USP 42,(62)	Absent/Present

END

Quality Manager