DAFFOHILS LABORATORIES PVT. LTD.

A WHO-GMP, GLP& ISO CERTIFIED COMPANY
F 109-110, UPSIDC INDUSTRIAL AREA, SELAQUI, DEHRADUN (U.K)-248011

QUALITY CONTROL DEPARTMENT

FINISHED PRODUCT CERTIFICATE OF ANALYSIS

Product Name	**ASTHANOX SYRUP**		
Generic Name	Salbutamol Sulphate Syrup IP		
Batch No.	DLS-26016	**A.R. No.**	DLPL/QC/FG/26/0077
Mfg. Date	09/2026	**Exp. Date**	08/2028
Batch Size	20,000 Bottles	**Sampled Qty**	12 x 100 ml
Date of Sampling	28/09/2026	**Date of Release**	29/09/2026

Sr. No.	TEST/PARAMETER	OBSERVED RESULTS	SPECIFICATION LIMIT
1	**Description**	Pink coloured liquid filled in amber colour round PET bottle.	Pink coloured liquid filled in amber colour round PET bottle.
2	**Identification (By HPLC)**	In the assay, the retention time of principal peak in the chromatogram obtained with the test solution is corresponded to the peak in the chromatogram obtained with the reference solution.	In the assay, the retention time of principal peak in the chromatogram obtained with the test solution is corresponded to the peak in the chromatogram obtained with the reference solution.
3	**pH**	4.23	3.4 to 4.5
4	**Nominal Volume**	100.14 ml	100.0 ml
5	**Weight per ml**	1.049 g/ml	1.0 g/ml to 1.40 g/ml
6	**Related Substance**		
	Impurities D	Not Detected	NMT 0.6 %
	Impurities F	Not Detected	NMT 0.6 %
	Impurities N	Not Detected	NMT 0.3 %
	Impurities J	Not Detected	NMT 0.3 %
	Impurities O	Not Detected	NMT 0.3 %
	Impurities Q	Not Detected	NMT 0.3 %
	Any Other secondary Peaks	0.05 %	NMT 0.2 %
	Total secondary Impurities	0.05 %	NMT 2.0%

	Prepared By	Checked By	Approved By
Name & Designation	**Officer Quality Control**	**Executive Quality Control**	**Manager Quality Control**
Signature			APPROVED
Date	29/09/26	29/09/2026	29/09/2026

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7	**Assay:**		
	Each 5ml contains:		
	Salbutamol Sulphate IP Eq. to Salbutamol 2 mg	2.0718 mg 103.59 %	1.9 mg to 2.1 mg 95.0% to 105.0%
8	**Additional Test**		
	Ethylene Glycol	Not detected	NMT 0.1%
	Diethylene Glycol	Not detected	NMT 0.1%

Remarks: The product complies/~~does not c~~omply with the above In-~~House~~/IP/~~BP/USP/JP~~ specification.

	Prepared By	**Checked By**	**Approved By**
Name & Designation	**Officer Quality Control**	**Executive Quality Control**	**Manager Quality Control**
Signature			APPROVED
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