

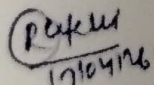
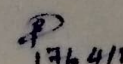
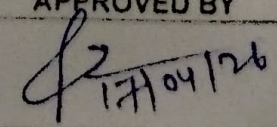


**M/S IVM PHARMACIA**  
Plot No. 05, Industrial Township,  
Bhatoli- Kalan, Baddi (H.P.) -173205  
**CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)**

**QUALITY CONTROL DEPARTMENT**

PRODUCT NAME	Levosalbutamol Hydrochloride and Budesonide Inhalation Suspension	BATCH NO.	12126002
GENERIC NAME	Levosalbutamol Hydrochloride and Budesonide Inhalation Suspension	A. R. NO.	QC/FP/26/0514
MFG. DATE	03/2026	PRODUCT CODE	21
EXP. DATE	02/2028	BATCH SIZE	300 LTR
SAMPLE SIZE	140 X 2 ml	PACK SIZE	2 ML
DATE TEST STARTED	02/04/2026	DATE TEST COMPLETED	----
MFG. LIC. NO.	MB/18/1020	SPECIFICATION NO.	FPS/F059-00

S.NO.	TEST	OBSERVATION	SPECIFICATION
1	Description	A white homogenous redispersible suspension.	A white homogenous redispersible suspension.
2	Identification (By HPLC) A) For Levosalbutamol Hydrochloride  B) For Budesonide	Complies  Complies	The retention time of major peak in Sample preparation solution is concordant with standard preparation solution.  The retention time of major peak in Sample preparation solution is concordant with standard preparation solution.
3	pH	4.58	3.5 to 5.0
4	Extractable Volume	2.1 ml	Not less than 2.0 ml
5	Sterility	Under test	There should be no observed microbial growth within 18 days.
6	Uniformity of Content	<b>For Levosalbutamol Hydrochloride</b> Average = UT Minimum = UT Maximum = UT  <b>For Budesonide</b> Average = UT Minimum = UT Maximum = UT	The preparation complies with the test if. each individual content is 85 to 115 per cent of the average content. The preparation fails to comply with the test if more than one individual content is outside these limits or if one individual content is outside, the limits of 75 to 125 per cent of the average content. If one individual content is outside the limits of 85 to 115 per cent of the

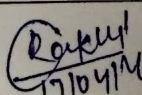
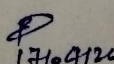
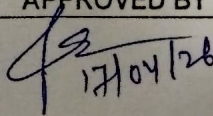
	ANALYSED BY	CHECKED BY	APPROVED BY
Signature/ Date	 17/04/26	 17/04/26	 17/04/26
Name	Rakul Baly	Bantu Kymon	
Designation	Officer-QC	Executive-QC	Quality Head

**M/S IVM PHARMACIA**Plot No. 05, Industrial Township,  
Bhatoli- Kalan, Baddi (H.P.) -173205**CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)****QUALITY CONTROL DEPARTMENT**

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			average content but within the limits of 75 to 125 per cent, repeat the determination using another 20 dosage units. The preparation complies with the test if not more than one of the individual contents of the total sample of 30 dosage units is outside 85 to 115 per cent of the average content and none is outside the limits of 75 to 125 per cent of the average content.
7	Assay (By HPLC): Each 2.0 ml respule contains: Levosalbutamol Hydrochloride IP ..... 1.25mg  Budesonide IP..... 0.5mg	1.265mg/2 ml (101.23%)  0.506 mg/2 ml (101.26%)	1.125mg/2ml to 1.375mg/2ml (90.0% and 110.0%)  0.463mg/2ml to 0.550mg/2ml (92.5% and 110.0%)

**Remark:** The above sample complies / does not comply as per IP/ BP/ USP specification or external lab analysis report.

	ANALYSED BY	CHECKED BY	APPROVED BY
Signature/ Date	 17/04/26	 17/04/26	 17/04/26
Name	Rakhi Bala	Anurag Kumar	Ramesh Singh
Designation	Officer-QC	Executive-QC	Quality Head