

METAPHARMACEUTICAL

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AVIK PHARMACEUTICAL LIMITED

A- 1/7 & A- 1/8, 1st Phase, GIDC, Vapi - 396 195, Dist-Valsad, Gujarat State, India

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CERTIFICATE OF ANALYSIS

Name of Product : BETAMETHASONE VALERATE EP		CAS. No : [2152-44-5]
IUPC name : 9-Fluoro-11 β , 21-dihydroxy-16 β -methyl-3, 20-dioxopregna-1, 4-diene-17-yl pentanoate.		
Batch No. : BV/M/007/25	A. R. No. : AVK/BV/011/25	
Mfg. Date : Mar.- 2025	Exp. Date : Feb.- 2030	
Batch size : 24650 gm	Batch out put : 24280 gm	
Date of released : 01/05/2025		

TESTS	SPECIFICATIONS	RESULTS
CHARACTERS		
APPEARANCE	White or almost white crystalline powder.	White crystalline powder.
SOLUBILITY	Practically insoluble in water, freely soluble in acetone and in Methylene chloride, soluble in ethanol (96 %).	Conforms
MELTING POINT	About 192 °C, with decomposition.	192.4°C
IDENTIFICATION	<p>A) By I.R: Infrared absorption spectrum of the sample should be in concordant with the IR spectrum of Betamethasone 17-Valerate working standard.</p> <p>B) By TLC: The principal spot in the chromatogram obtained with the test solution is similar in position, colour, and size to the principal spot in the chromatogram obtained with the reference solution</p> <p>C) By HPLC: The principal peak in the chromatogram obtained with the test solution is similar in retention time and size to the principal peak in the chromatogram obtained with reference solution (b).</p> <p>D) By chemical: The colour is discharged and a clear solution remains</p>	<p>Conforms</p> <p>Conforms</p> <p>Conforms</p> <p>Conforms</p>
SPECIFIC OPTICAL ROTATION	+ 77° to + 83° (dried substance).	+81°
RELATED SUBSTANCES (BY HPLC)	<p>Impurity A : NMT 0.7 %</p> <p>Impurity B : NMT 0.15 %</p> <p>Impurity I : NMT 0.15 %</p> <p>Impurity C : NMT 0.15 %</p> <p>Impurity H : NMT 0.15 %</p> <p>Impurity D : NMT 0.15 %</p> <p>Impurity E : NMT 0.3 %</p> <p>Impurity G : NMT 0.3 %</p> <p>Unspecified impurity : NMT 0.10 %</p> <p>Total impurities : NMT 1.5 %</p>	<p>ND</p> <p>ND</p> <p>ND</p> <p>ND</p> <p>ND</p> <p>ND</p> <p>0.23 %</p> <p>ND</p> <p>0.04 %</p> <p>0.27 %</p>
LOSS ON DRYING	Maximum 0.5 %	0.3 %
ASSAY (By UV)	Not Less Than 97.0 % - 103.0 % (Dried basis)	99.2 %
A) RESIDUAL SOLVENTS	<p>Methanol : NMT 3000 ppm</p> <p>Acetone : NMT 5000 ppm</p> <p>Methylene chloride : NMT 600 ppm</p> <p>Ethyl acetate : NMT 5000 ppm</p>	<p>Below detection limit</p> <p>907 ppm</p> <p>Below detection limit</p> <p>84 ppm</p>
B) PARTICLE SIZE (BY VOLUME BASIS)	99 % < 20 μ m	7.10 μ m

REMARKS: The Batch CONFORMS as per EP-11.4 Specifications.

<p>PREPARED BY:</p> <p>NAME: PRATIK PATEL</p> <p>DESIGNATION: EXECUTIVE-QC</p> <p>DATE: 01/05/2025</p>	<p>CHECKED BY:</p> <p>NAME: HEMANT KEVAT</p> <p>DESIGNATION: MANAGER-QC</p> <p>DATE: 01/05/2025</p>	<p>APPROVED BY:</p> <p>NAME: VINU PATEL</p> <p>DESIGNATION: SR. MANAGER-QA (DOC)</p> <p>DATE: 01/05/2025</p>
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