

ALVENTA PHARMA LIMITED VILL. KISHANPURA, TEHSIL BADDI- NALAGARH ROAD, DISTT.- SOLAN (H.P) 174101

Quality Control Department

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

Product Name	Vildafox-50	A.R. No.	FG/G/24A0166
Generic Name	Vildagliptin Tablets IP 50 mg	Sample Quantity	60 Tablets
Batch No.:	AGT40276	Sample Received on	04/03/2024
Batch Size:	1.70 Lac	Analysis Date	04/03/2024
Mfg. Date.	02/2024	Release Date	11/03/2024
Exp. Date	01/2027	Page No.:	Page 1 of 1

Sr. No.	Test Parameter		Acceptance Criteria Result					
1.	Description	17				colour, round shaped, biconvex de plain, uncoated tablet.		
2.	Identification	chron soluti chron	natogram o	principal peak in the btained with the test ands to the peak in the obtained with the in (a).	Complies			
3.	Average weight	180.0	$0 \text{ mg} \pm 3.0\%$	6. ·	181.86 mg			
4.	Uniformity of weight	± 7.5°	% of its ave	rage weight.	Min: 178.86 mg ; Max: 183.42 mg -1.65% ; +0.86%			
5.	Disintegration	Not n	nore than 15	minutes.	04 minutes 12 seconds.			
6.	Friability	Not n	nore than 1.0)%w/w	0.32 %w/w			
7.	Hardness	Not le	ess than 4.0	Kg/cm ²	5.52 Kg/cm ²			
8.	Dissolution	1		00% (Q) of labeled, d in 30 minutes.	Minimum = 97.40% Maximum = 100.22% Average = 99.15%			
9.	Assay:							
	Each uncoated tablet contains:		Claim	Limit		mg	%	
	Vildagliptin IP		50 mg	Between 95.0 % to 105.0 % of abeled amount. (Between 47.5 mg to 52.5 mg)		51.33 mg	102.65%	
10.	Microbial Limit Tests:		3					
i.	Total aerobic microbial count NMT 1000 cfu/g				20 cfu/g			
ii.	Total yeast and mould count NMT 100 cfu/g			cfu/g	Less than 10 cfu/g			
iii.	Pathogens: Escherichia col	i	Should be	absent/g		Absent		

Remarks: The above test parameters are complies/ not complies as per IP/BP/USP & In-House Specification.

Particulars	Prepared By	Checked By	Approved By
Name	Braveen Kynnor	DURGESHKUMAR	Co la sinal
Designation	Fre I	AC Head	Quy gan singh
Signature	Putre		Hend-Gilt
Date	11/03/2-24	11/03/2024	11/03/2024