

QUALITY CONTROL DEPARTMENT

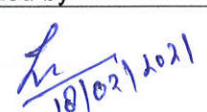
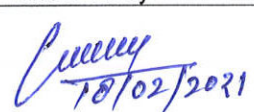
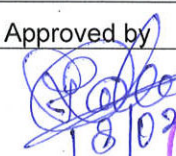
**CERTIFICATE OF ANALYSIS
(FINISHED PRODUCT)**

(Under Drugs &Cosmetics Act 1940 and Rules Made There Under)

Product Name	Dutacap	A.R. No.	: CF21-0153
Generic Name	Dutasteride Capsules IP 0.5 mg		
Batch No.	AC21041	Mfg. Date	: FEB.2021
Batch Size	1.0 Lac	Exp. Date	: JAN.2023
Pack size	10 X 10 Capsules	Receipt Date	: 17/02/2021
Sample Qty.	60 Capsules	Release Date	: 18/02/2021
Manufactured for:	Naiom Healthcare		
Manufacturing Lic. No.	82/UA/2006		

S.N.	Tests	Specification	Results	
1.	Description	Dark brown colour cap and body hard gelatin capsule fill white colour powder.	Dark brown colour cap and body hard gelatin capsule fill white colour powder.	
2.	Identification	In the Assay, the retention time of principal peak in the chromatogram obtained with the test solution corresponds to the peak in the chromatogram obtained with the reference solution.	Complies	
3.	Related substances	Singal Impurity NMT 1.0%	Not detected	
		Total Impurity NMT 2.0%	Not detected	
4.	Avg. weight of capsules	496 mg ± 7.5%	495.36 mg	
5.	Net Content	400 mg ± 7.5%	401.12 mg	
6.	Uniformity of weight	± 7.5% of the average weight	Within limit	
7.	Dissolution	D. Not less than 80 per cent	89.63 % to 97.56%	
8.	ASSAY:			
	Each hard gelatin capsule contains	Claim	Result	limit
	Dutasteride IP	0.5 mg	0.496 mg (99.20%)	NLT 90% - NMT 110%

Opinion: in the opinion of the undersigned, the sample referred to above is of **STANDARD QUALITY** as per In-House Specification.

Analyzed by	Checked by:	Approved by
 Sign/Date: Officer QC	 Sign/Date: Executive QC	 Sign/Date: Manager QC

