

Pace Biotech

Surajpur, Paonta sahib Dist. Sirmour (H.P)

Title :

Certificate of Analysis Finished Product

Product Name	ESOMECAN 40 Injection	A.R. No.	NB/FP/24/084
Generic Name	Esomeprazole Sodium for Injection 40mg	Sampled qty.	45 Vials
Batch No.	N24070F	Sampled by	Aakash
Batch Size	15,000 Vials	Sampled on	25/06/2024
Mfg. Date	06/2024	Date of Testing	25/06/2024
Exp. Date	05/2026	Date of Release	10/07/2024

S. No.	Tests	Specifications		Observations		
1.	Description	A white dry powder filled in clear colour glass vial.			A white dry powder filled in clear colour glass vial.	
2.	Identification (By HPLC)	In the chromatogram obtained the retention time of major peak in test solution is correspondence with RT of major peak in working standard solution.			Complies	
3.	Uniformity of weight	Average weight ± 10%		-3.60% & +2.87%		
4.	Average weight	Informative.		120.4 mg		
5.	Particulate Matter (a.) Sub-Visible particle count (1.) Particles ≥10µm (2.) Particles ≥25µm (b.) Visual	NMT-6000/vial NMT-600/vial The solution should be essentially free from extraneous, mobile undissolved particles, other than gas bubbles, unintentionally.		1382/vial 8/vial Complies		
6.	Water (By K.F)	NMT- 5% w/w		4.06% w/w		
7.	Sterility	No microbial growth should be Complies observed.				
8.	Bacterial Endotoxins	NMT 5.0 EU/mg of Esomeprazole. Less than Esomepra			5.0 EU/mg of zole.	
9.	Assay: Each vial Contains					
Ingredients		Labeled Claim	Found		of labeled amount	Limits % of labeled amount
Esomeprazole Sodium (Sterile) Eq. to Esomeprazole (suitably buffered) (Lyophilized)		40 mg	38.53 mg		96.33%	90.0 to 110.0%

Remarks: In the opinion of the undersigned the sample referred to above is/ is not of the standard quality as The Drug & Cosmetic Act 1940 and the rules made there under. Complies/not complies as per IP/BP/USP/IHS.

Analysis by

Checked by

Approped by