



D.R. JOHN'S LAB (P) LTD.
Plot No. 3, Sector 6-A, IIE, SIDCUL, Haridwar-249403 (Uttarakhand)

QUALITY CONTROL DEPARTMENT

**CERTIFICATE OF ANALYSIS
(FINISHED PRODUCT)**

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

Product Name	Rabezcan-DSR Capsules	A.R. No.	:DRJF-250221
Generic Name	Rabepazole Sodium (EC) & Domperidone (SR) Capsules		
Batch No.	CTP-25032	Mfg. Date	: Jul.2025
Batch Size	200000 cap	Exp. Date	: Jun.2027
Pack size	10 X 10 Capsules	Receipt Date	: 02-07-2025
Sample Qty.	50 Capsules	Release Date	: 05-07-2025
Manufactured for	American Remedies Healthcare Pvt.		
Manufacturing Lic. No.	113/UA/2007		

Sr.No.	Tests	Specification	Results
1.	Description	Transparent and pink colour elongated shape hard gelatin capsule containing coloured pellets.	Transparent and pink colour elongated shape hard gelatin capsule containing coloured pellets.
2.	Identification	In the Assay, the principal peak in the chromatogram obtained with the test solution corresponds to the chromatogram obtained with the reference solution.	Complies
3.	Ave. weight of capsules	350.0 mg \pm 7.5 %	360.54 mg
4.	Net Content	300 mg \pm 7.5 %	296.83 mg
5.	Uniformity of weight	\pm 7.5 % of average weight	-3.07% to +3.08%
6.	Dissolution (Domperidone) IN 1 hour IN 4 hour IN 8 hour IN 12 hour	(15.0% to 40%) (30.0% to 60%) (55.0% to 85%) (NLT-70%)	25.13% 51.06% 75.59% 89.10%
7.	Dissolution Rabepazole Sodium In 0.1N HCl	Not less than 85% of labeled amount of Rabepazole shall retain as residue after 2 hours	97.28 %
8.	In Buffer	(Q) NLT - 70.0 %	88.63 %
9.	Assay.		

Each hard gelatin capsule contains:	Claim	Limit	Result
Rabepazole Sodium IP As enteric coated pellets	20 mg	NLT 90% TO NMT 110%	19.83 mg (99.15%)
Domperidone IP As sustained release pellets	30 mg	NLT 90% TO NMT 110%	30.35 mg (101.17%)

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

Analyzed by <i>Devendra</i> 05/07/25 Sign/Date: Officer QC	Checked by: <i>U.S.</i> 05/07/25 Sign/Date: Executive QC	Approved by <i>U.S.</i> 05/07/25 Sign/Date: Manager QC
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