

Plot No. A-67 to 72, PIPDIC Electronic Park, Thirubuvanai, Puducherry-605107 Tel: 0413 – 2641099 2641199 2641666 info@safetab.net www.safetab.net

CERTIFICATE OF ANALYSIS						
Product Name	:	LITACOLD FLU				
		(Paracetamol, Phenylephrine,	(Paracetamol, Phenylephrine, Chlorphenamine Maleate & Caffeine Tablets)			
Batch No.	:	G17250716 Batch Size : 12.5LAC				
Mfg. Date	:	JUL'2025	Exp. Date	:	JUN'2028	
Sampling Date	:	25/07/2025	Sample Quantity	:	13X4'S	
Analysis Date	:	25/07/2025				
Release Date	:	30/07/2025	A.R. No.	:	SFP251231	

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Sr. No.	Test	Specifications	Results		
1.	Description	Light yellow coloured, flat, round beveled edged uncoated tablet with break line on one side and plain on other side.	Light yellow coloured, flat, round beveled edged uncoated tablet with break line on one side and plain on other side.		
2.	Identification (By HPLC)				
	A. Paracetamol BP	The retention time of one of major peak in the chromatogram of the sample preparation corresponds to the peak due to Paracetamol in the standard preparation as obtained in assay.	Complies		
	B. Phenylephrine HCI BP	The retention time of one of major peak in the chromatogram of the sample preparation corresponds to the peak due to Phenylephrine HCl in the standard preparation as obtained in assay.	Complies		
	C. Chlorphenamine Maleate BP	The retention time of one of major peak in the chromatogram of the sample preparation corresponds to the peak due to Chlorphenamine Maleate in the standard preparation as obtained in assay.	Complies		
	D. Caffeine BP	The retention time of one of major peak in the chromatogram of the sample preparation corresponds to the peak due to Caffeine in the standard preparation as obtained in assay.	Complies		
3.	Average Weight of tablets	635.0 mg ± 3.0% (615.9 mg to 654.0 mg)	635.0mg		

Particulars	Prepared By	Reviewed By	Approved By
Name	C.K.SARAVANAN	K.SARAVANAN	M.VIJAYAKUMAR
Designation	Asst. Manager - QC	Dy. Manager - QC	GM - QC
Signature/Date			

Format No: ST/QC/070:F7



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Sr. No.	Test	Specifications	Results		
4.	Uniformity of weight	Not more than 2 of the individual weights deviate from the average weight by more than ± 5% and none deviate by more than ± 10.0%	(-) 1.51% (+) 1.23%		
5.	Diameter	12.70 mm ± 0.2 mm (12.50 to 12.90 mm)	12.79mm		
6.	Thickness	4.30 mm ± 0.2 mm (4.10 to 4.50 mm)	4.17mm		
7.	Hardness	100 N to 250 N	186.95N		
8.	Friability	Not more than 1.00%	0.17%		
9.	Disintegration time	Not more than 15 minutes	05 minutes 31 seconds		
10.	Dissolution:				
	Chlorphenamine Maleate BP 2 mg	Not less than 80.0% of the labeled amount	Min: 98.5%; Max: 100.4%; Avg: 99.5%		
	Phenylephrine HCI BP5 mg	Not less than 80.0% of the labeled amount	Min: 98.2%; Max: 99.7%; Avg: 98.9%		
	Caffeine (anhydrous) BP30 mg	Not less than 80.0% of the labeled amount	Min: 99.4%; Max: 101.4%; Avg: 100.5%		
	Paracetamol BP 500 mg	Not less than 80.0% of the labeled amount	Min: 100.1%; Max: 101.5%; Avg: 100.9%		
11.	Uniformity Content:				
	Chlorphenamine Maleate BP 2 mg	Not less than 85.0 % and not more than 115.0 % of labeled claim	Min: 97.9%; Max: 104.0%; Avg: 100.7%		
	Phenylephrine HCI BP5mg	Not less than 85.0 % and not more than 115.0 % of labeled claim	Min: 97.9%; Max: 104.0%; Avg: 100.4%		

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Test	Specifications	Results					
Related Substances							
Single maximum unknown impurity	Not more than 0.20%	Not Detected					
Total impurities	Not more than 0.50%	Not Detected					
Assay: Each uncoated ta	blet contains:						
Chlorphenamine Maleate BP 2 mg	1.9mg to 2.2mg (95.0% to 110.0% of the labeled claim)	2.01mg (100.6%)					
Phenylephrine HCI BP5 mg	Not less than 95.0% and not more than 110.0% of labeled claim	4.94mg (98.9%)					
Caffeine (anhydrous) BP30 mg	Not less than 95.0% and not more than 110.0% of labeled claim	30.93mg (103.1%)					
Paracetamol BP 500mg	Not less than 95.0% and not more than 110.0% of labeled claim	498.89mg (99.8%)					
Microbial Limits:							
Total aerobic bacterial counts	Not more than 1000 cfu/g	10cfu/g					
Fungi	Not more than 100 cfu/g	<10cfu/g					
E. coli.	Should be Absent	Absent					
Salmonella.	Should be Absent	Absent					
S. aureus.	Should be Absent	Absent					
P. aeruginosa. Should be Absent		Absent					
	Related Substances Single maximum unknown impurity Total impurities Assay: Each uncoated ta Chlorphenamine Maleate BP	Related Substances Single maximum unknown impurity Total impurities Not more than 0.20% Assay: Each uncoated tablet contains: Chlorphenamine Maleate BP					

Remarks: The sample Complies/Does not Complies as per BP / USP / In-House specification.

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