

M.L.A. Industries Pvt. Ltd. (Pharma Division)

111/230,HARSH NAGAR, KANPUR, Tel.: (0512)-2555292 Website: www.mlagroup.com, E-mail:sales@mlagroup.com

(Drug License No. UP78250000012)

(Food Safety & Drug Administration - Govt. of Uttar Pradesh)

CERTIFICATE OF ANALYSIS

CUSTOMER: M/s

PRODUCT: CALCIUM STEARATE

BATCH NO.: CS-21C012023

Date of MFG.: 21/01/2023

INVOICE: Sample

GRADE: IP

DATE: 11/02/2023

Date of EXP: 5 years from MFG

PROPERTY	UNIT	SPECIFICATIONS	TEST VALUE
Description	-	White Free Flowing Powder	White Free Flowing Powder
Purity (assay)	%	9.0 - 10.5	9.78
L.O.D. (105°C)	%	4.0 Max	2.50
Identification	-	Test A –The H2O layer gives the test for Calcium	Complies to IP
		Test B- Passes, Fatty acid obtained not congeal below 54 °C	
Heavy Metals as Lead	ppm max	10	< 10
Composition of Fatty Acid	%	Percentage of stearic acid in fatty acid NLT-40.0	
		& Percentage sum of stearic acid and palmic acid NLT-90	Complies to IP

Tested by Chemist	Checked by QA	Approved by Lab Head
Engura 9-	P. Wmar 25/02/23	UNIVO 212013 RESPHARMAD
Mr. Gaurav Sen	Mr. Prince Srivastava	Mr. Ujjval Tiwari



RIGHT CHEMISTRY, RIGHT PEOPLE



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Regd. Office: 111/230, HARSH NAGAR, KANPUR, Tel.: (0512)-2555292 Factory Address: D-7B, D-8 & D-16, Panki Industrial Area, Site-1, Kanpur, Uttar Pradesh 208022 Website:www.mlagroup.com, E-mail:sales@mlagroup.com

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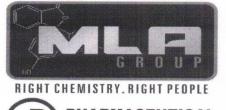
CERTIFICATE OF ANALYSIS

CUSTOMER: M/s : Sample **INVOICE** PRODUCT: CALCIUM STEARATE : BP GRADE BATCH NO.: CS-22B012021 : 28/02/2023 DATE

Date of MFG: 22/01/2023 Date of EXP: 5 years from MFG

PROPERTY	UNIT	SPECIFICATIONS	TEST VALUE
Colour & Appearance	-	White Free Flowing Powder	Complies with BP
Solubility	-	Practically insoluble in water and in ethanol (96 percent).	Complies with BP
Acidity Or Alkalinity	-	NMT 0.5 ml of 0.01 M Hcl or 0.01 M Naoh Required For Change the colour.	Complies with BP
Identification	-	Test A – Freezing Point-53°C Min Test B- Acid Value between 195-210 Test C – Passes Test D - Passes	A - 59 °C B - 204 C – Complies with BP D - Complies with BP
Acidity Or Alkalinity	-	NMT 0.5 ml of 0.01 M Hcl or 0.01 M Naoh Required For Change the colour.	Complies with BP
Chloride	% max	0.1	<0.1
Sulphates	%max	0.3	<0.3
Cadmium	ppm max	3.0	0.22
Lead	ppm max	10.0	3.1
Nickel	ppm max	5.0	0.72
Loss on Drying	% Max.	6.0 % Max.	3.14
Microbial Contamination	CFU/g	TAMC-NMT 10 ³ ,TYMC-NMT 10 ² , Salmonella & Escherichia Coli-Absent	Complies with BP
Assay As Ca	%	6.4 - 7.4	7.18
Fatty Acid Composition			
1. Stearic Acid	%	For 1. NLT 40%	Complies with BP
2. Sum of Palmitic acid	%	For 2. NLT 90%	1
+St.Acid		NLT=Not Less Than	
Particle Size Distribution	-	Passes Complies w	
Specific Surface Area	M ² /g	3.0 - 14.0	10.28

Tested by Chemist	Checked by QA	Approved by Lab Head	
28/02/23	P. Komaro	WW 23 23 RESPHARMA OIL	
Mr. Gaurav Sen	Mr. Prince Srivastava	Mr. Ujjval Tiwari	
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CERTIFICATE OF ANALYSIS

CUSTOMER:	INVOICE : Sample
PRODUCT : CALCIUM STEARATE	GRADE: USP
BATCH NO.: CS-18A022023	DATE :28/02/2023
Date of MFG: 18/02/2023	Date of EXP: 5 years from MFG

UNIT	SPECIFICATIONS	TEST VALUE
-	White Free Flowing Powder	White Free Flowing Powder
-	Test A – Freezing Point- 53°C Min	Test A –57
	Test B- Fatty acid obtained not congeal below 54 °C	Test B- Passes
0/0	Ca- 6.4 - 7.4	6.85
%	NLT 40 %	Passes
%	NLT 90 %	Passes
ppm max	10	< 10 ppm
%	NMT 4.0 @105 *C temp	3.5
	% % ppm max	- White Free Flowing Powder - Test A – Freezing Point- 53°C Min Test B- Fatty acid obtained not congeal below 54°C % Ca- 6.4 - 7.4 % NLT 40 % NLT 90 % ppm max 10

Tested by Chemist	Checked by QA	Approved by Lab Head	
Layras - 28/02/23	P. hemas. 28/02/23	UNITED TO THE PHARMA DILLEGE	
Mr. Gaurav Sen	Mr. Prince Srivastava	Mr. Ujjval Tiwari	
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RIGHT CHEMISTRY. RIGHT PEOPLE



Date of MFG.: 16/02/2023

M.L.A. Industries Pvt. Ltd. (Pharma Division)

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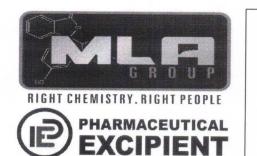
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CERTIFICATE OF ANALYSIS

INVOICE : Sample **CUSTOMER: M/s** : EP : CALCIUM STERATE **GRADE PRODUCT** : 01/03/2023 DATE BATCH NO.: CS-16E022023 Date of EXP: 5 years from MFG

PROPERTY	UNIT	SPECIFICATIONS	TEST VALUE
Description	-	Appearance: fine, white or almost white, crystalline powder.	Complies to EP
Solubility		practically insoluble in water and in ethanol.	Complies to EP
Identification	A. Freezing point : minimum 53 °C		A – 58 °C B - 204 C - Complies to EP D - Complies to EP
Acidity or alkalinity	-	Not more than 0.5 ml of 0.01 m hcl or 0.01 M sodium hydroxide is required to change the colour of the indicator.	Passes
Chloride	%	NMT 0.1	< 0.1
Sulfates	%	NMT 0.3	<0.3
Cadmium	ppm	NMT 3.0	0.34
Lead	ppm	NMT 10.0	4.89
Nickel	ppm	NMT 5.0	0.61
Loss on drying	%	NMT 6.	2.84
Microbial contamination	CFU/g	TAMC: acceptance criterion 10 ³ CFU/g TYMC: acceptance criterion 10 ² CFU/g E.coli & Salmonella- Absent	Passes
Assay (Calcium)	%	6.4 - 7.4	7.06
Fatty Acid Composition 1. Stearic Acid 2. Sum of Palmitic acid +St.Acid	%	For 1. NLT 40.0 For 2. NLT 90.0	Complies to EP
Particle Size	-	Passes	Complies to EP

Tested by Chemist	Checked by QA	Approved by Lab Head
C/103/23	P.Kumar 0 01/03/2023	UNINO 103123 RESPRANTADA
Mr. Gaurav Sen	Mr. Prince Srivastava	Mr. Ujjval Tiwari
		* KANPUR



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CERTIFICATE OF ANALYSIS

CUSTOMER:	INVOICE:	`
PRODUCT : CALCIUM STEARATE	GRADE: JP	-
BATCH NO.: 18E0202023	DATE: 01/03/2023	
Date of MFG. : 18/02/2023	Date of EXP: 5 years from MFG	

PROPERTY	UNIT	SPECIFICATIONS	TEST VALUE
Description	-	Calcium Stearate occurs as a white, light, bulky powder. It feels smooth when touched, and is ladhesive to the skin. It is odorless or has a faint, characteristic odor.	Complies to JP
SOLUBILITY	-	It is practically insoluble in water, in ethanol (95) and in diethyl ether.	Complies to JP
Identification	-	Test A – Shake vigorously 3g Stearate with 20 ml. of diluted hydrochloric acid (1 in 2) and 30 mL of diethyl ether for 3 minutes, and allow to stand: the separated aqueous layer responds to the Qualitative Tests (1), (2) and (4) for calcium salt.	Passes
		Test B – residue melts at a temperature not below 54*C.	Passes
Heavy metals	ppm	NMT 20 ppm .	Passes
Arsenic	ppm	NMT 2 ppm.	Passes
Loss on drying	%	NMT 4.0% (1 g, 105*C, 3 hours)	2.85
Assay	%	NLT 6.4% and NMT 7.1% of calcium	6.86

Tested by Chemist	Checked by QA	Approved by Lab Head
hayrav	P. h. maso 01 (23/23	JAN DO 19323 REES PHARMADILE
Mr. Gaurav Sen	Mr. Prince Srivastava	Mr. Ujjval Tiwari
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