

	CERTIFICAT DE CONFORMITE	DATE
		03/06/19
		VERSION N°1

POUR LE CLIENT

NUV

NOVOMA SARL
 BATIMENT ZEPHYR AVENUE BERNARD
 31 400 TOULOUSE

Je soussigné M. WACRENIER, Président de Laboratoire PHYTOCOSMA SAS certifie que le produit cité ci-après est conforme aux spécifications établies.

Nous vous rappelons qu'il vous appartient de vérifier les conditions de distribution et d'utilisation de ces produits conformément à la législation en vigueur.

Code interne	NUVHAY02
Désignation interne	ACIDE HYALURONIQUE

Code client	3 770025 711324
Désignation client	ACIDE HYALURONIQUE + Vitamine C

Numéro de lot	D16838	Numéro de BL	20881
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Date de fabrication	17/10/2023	DDM	10/2026
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Conditions de conservation	A conserver à l'abri de l'oxygène et de la lumière à une température comprise entre 15 et 25°C dans son emballage d'origine
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Le produit contient de(s) Allergène(s)	Non
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Liste des allergène(s) dans le produit

Non applicable

Le produit contient de(s) Additif(s)	Non
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Liste de(s) additif(s) dans le produit

Non applicable

Le produit est BIO	Non
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(*Produit issu de l'agriculture biologique FR-BIO-01 et process conforme à la fabrication de produits biologiques)

Le produit est sans OGM	Oui	Le produit est Ionisé	Non
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Le produit est sans Gluten	Non
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Conforme Végétarien	Oui	Conforme Végétalien	Oui
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Conforme Halal	Oui	Conforme Casher	Oui
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CERTIFICATE OF ANALYSIS

ExceptionHYAL[®] Star

Batch n.	23L077/S
Production date	JULY 2023
Period of best use	JULY 2026

Parameter	Unit	Requirement	Method	Result
PHYSICAL-CHEMICAL				
Appearance	-	Powder	IO 07-06A	Powder
Colour	-	White	IO 07-06A	White
Odour	-	Characteristic	IO 07-06A	Characteristic
pH (0,1% solution)	-	6,0 – 8,0	IO 07-06B	6,0
Transparency (0,1% solution,550 nm)	%	≥99	Ph.Eur. 2.2.25	Complies
Loss on drying (105°C)	%	≤10	IO 07-06K	5
Ashes	%	≤15	Ph.Eur. 2.4.16	Complies
Bulk density	g/cm3	0,3 -0,5	Ph.Eur. 2.9.34	0,4
Glucuronic acid assay	%	≥46	S.H. Ph.Eur.Monograph	95
Sodium Hyaluronate assay	%	≥95	S.H. Ph.Eur.Monograph	46
Lead (Pb)	ppm	≤3	IO 07-06AG	Complies
Cadmium (Cd)	ppm	≤1	IO 07-06AG	Complies
Mercury (Hg)	ppm	≤0,1	IO 07-06AG	Complies
Water activity	-	≤0,5	IO 07-06AL	0,1
MICROBIOLOGICAL				

CERTIFICATE OF ANALYSIS

ExceptionHYAL[®] Star

Total bacterial count	CFU/g	≤100*	Ph.Eur. 2.6.12/USP (current edition)	<100
Yeasts & Moulds	CFU/g	≤100*	Ph.Eur. 2.6.12/USP (current edition)	<100
<i>Enterobacteriaceae</i>	CFU/g	≤100	UNI ISO 21528-2	<100
<i>Escherichia coli</i>	g	Absence	Ph.Eur. 2.6.13/USP (current edition)	Absence
<i>Staphylococcus aureus</i>	g	Absence	Ph.Eur. 2.6.13/USP (current edition)	Absence
<i>Salmonella</i>	10g	Absence	Ph.Eur. 2.6.13/USP (current edition)	Absence

* 2x (as per Ph.Eur.5.1.4)

Date: 11th September 2023

Roelmi HPC Technical Department

The analyses are performed on a representative sample and are referred to the product at the time of release. The information included in this Certificate of Analysis does not discharge the user from the control of the product before the use. Roelmi HPC SRL does not take liability for any damage due to improper use. This document refers to the latest version of the specification data sheet and it has been prepared by electronic processing not requiring signature and the traceability to the original signature is managed by internal quality assurance system.

VC Ascorbic Acid 100 mesh 95%



Coversheet for Certificate of Analysis

Productcode : 5015842
Lot No. : HPB2202089
Analysis No. : 75752335

CfC Number : 04917410
Articlecode : 5015842368
Sales Name : VC Ascorbic Acid 100 mesh 95%,25KG BAL
Manufacturing Date : 11-FEB-2022
Best Use Before Date : 10-FEB-2025
Delivery Note No. 8322177950
Destination : Germany
Customer Ref. No : P22-20045/002
Customer Article No:
Customer Name: MDC GmbH
Zum Reiherhorst
Stelle

Article desc. CoA
Comment

VC Ascorbic Acid 100 mesh 95%,25KG BAL

CERTIFICATE OF ANALYSIS

Productcode	: 5015842368	Batch Size	: 3,000KG
Lot No.	: HPB2202089	Manufacturing Date	: 11.2.2022
Analysis No.	: 75752335	Best use before	: 10.2.2025
Report Date	: 21.2.2022		

Test	Result	Limits/Specification	Dimension/Units
*Appearance	Correspond	White or almost white, crystal or crystalline powder	
Solubility	Correspond	Freely soluble in water, slightly soluble in ethanol, insoluble in ether or chloroform	
Identification A	Positive	Positive	
Identification B	Correspond	IR	
*Melting Point	191	190 - 192	°C
*Specific Optical Rotation	20.9	20.5 - 21.5	°
*Clarity and Color of Solution	Correspond	Clear, not more intensely than BY7	
*pH(2%, W/V)	2.6	2.4 - 2.8	
*pH(5%, W/V)	2.4	2.2 - 2.5	
Loss on Drying	< 0.1	≤ 0.1	%
Residue on Ignition	< 0.1	≤ 0.1	%
Arsenic(as As)	< 1.0	≤ 1.0	ppm
Heavy Metals (as Pb)	Correspond	≤ 0.0003%	
Lead	< 1.0	≤ 1.0	ppm
Mercury	< 0.5	≤ 0.5	ppm
Iron	< 2	≤ 2	ppm
Copper	< 3	≤ 3	ppm
Oxalic Acid	Correspond	≤ 0.2%	
Residual Solvents(as methanol)	< 250	≤ 250	ppm
*Assay	99.9	99.0 - 100.5	%
Total Plate Counts	< 1000	≤ 1000	CFU/g
Yeasts & Moulds	< 100	≤ 100	CFU/g
Escherichia.Coli	Correspond	Absence in 1g	
Salmonella	Correspond	Absence in 25g	
Staphylococcus Aureus	Correspond	Absence in 25g	
Impurity C	< 0.15	≤ 0.15	%
Impurity D	< 0.15	≤ 0.15	%
Unspecified impurities	< 0.10	≤ 0.10	%
Total Impurities other than C & D	< 0.2	≤ 0.2	%
*Particle Size(through us 100mesh)	96	≥ 95	%
Cadmium	< 0.5	≤ 0.5	ppm

Conclusion: The product with this batch number meets all requirements when tested according to the monograph of USP/FCC/BP/EP/E300 in current version.

Test items marked with * are based on routine batch analysis. Others are tested at regular intervals.

This is an electronic COA. The data from batch analysis are in accord with the actual test data, which are true, correct and

VC Ascorbic Acid 100 mesh 95%,25KG BAL

CERTIFICATE OF ANALYSIS

Productcode	:	5015842368	Batch Size	:	3,000KG
Lot No.	:	HPB2202089	Manufacturing Date	:	11.2.2022
Analysis No.	:	75752335	Best use before	:	10.2.2025
Report Date	:	21.2.2022			

controlled. The data from periodic analysis are controlled at regular intervals as part of our assurance program.

QC Manager



Nie Xiaoming (Daniel Nie)

Quality Manager



Ge Liang

Certificate of Analysis

General Information

Product	Remy O DR6	Production Date	12/02/2023	(dd/mm/yyyy)
Batch	2320302850	Best before	11/02/2027	(dd/mm/yyyy)
Issued by	Quality Control Management	Date CoA Issued	08/03/2023	(dd/mm/yyyy)

Results of analyses

Parameter	Result	Unit	Method ⁽¹⁾	LSL	USL
Physical and Chemical Parameters					
Moisture	7	g/100g	ISO 712	≤	14
Protein content (N*6,25) on DM	3,6	g/100g d.m.	ISO 1871 ⁽¹⁾	≤	6,0
Ash content on DM	0,2	g/100g d.m.	ISO 3593	≤	1,0
Rheological Parameters					
Starting gel point, pH as is, 6%	83	°C	Brabender	≥	60
End viscosity, pH as is, 6%	584	BU	Brabender	≥	500
Microbiological Parameters					
Salmonella (/375g)	Negative	/375g	ISO 6579		Negative
Total mesophilic bacteria (aerobic)	100	cfu/g	ISO 4833	≤	100.000
Yeasts and Moulds	<10	cfu/g		≤	1.000
Enterobacteriaceae	<10	cfu/g		≤	100

⁽¹⁾ or (acknowledged and) validated equivalent

Remarks

We herewith confirm that the product complies with the corresponding guarantees listed in its Product Sheet .

Rice starch issued from organic farming, Certisys BE-BIO-01 , Certified Organic NOP by bio.inspecta AG

CERTIFICATE OF ANALYSIS EMPTY HARD CAPSULES OF VEGETABLE ORIGIN

CUSTOMER: GOCAPS GMBH			
LOT No.: K2209001976	PRODUCT CODE: K00016	SIZE: 1	
PURCHASE ORDER NUMBER: PO2000257	CHARGE No.: 1-000429	ART No.: 56-000108	
CAPSULE COLOR / CODE: CAP - NATURAL 1-0K / BODY - NATURAL 1-0K			
PRINT: N/A	TEXT: N/A	INK COLOR: N/A	

THIS IS TO CERTIFY THAT: The hard capsules of vegetable origin (K-CAPS) manufactured by C.I. FARMACAPSULAS S.A.S. are made from cellulose ethers, which are polymers derived from vegetable sources. Our capsules are certified as Kosher and Halal, and meet all requirements of current European Pharmacopoeia (EP) and United States Pharmacopoeia (USP). Hypromellose used in the manufacturing of capsules meet specifications as described in the current United States Pharmacopoeia. Cellulose ethers are considered as Generally Recognized As Safe (GRAS) by the FDA.

(% Ingredients to % Cellulose)

Cap	%	Body	%
<p><i>Colorant and ingredients used in capsules are officially approved for use as dye in Foods, Drugs and Cosmetics and/or Drugs and Cosmetics, in the country of destination.</i></p> <p><i>Some changes in color are due to natural colorants or can occur/are within the specification. The above specifications apply to all capsules having the same size and code numbers, unless otherwise stipulated.</i></p>			

Date of Manufacture: 2022-10


Expiration Date: 2027-10

CRITERIA	METHOD / REFERENCE	SPECIFICATIONS	RESULTS
PHYSICAL			
Average Capsule Weight	DCC-MI-P003/ USP<2091>	75.00-85.00 mg	78.1
Loss on drying	DCC-MA-P027	4.00-8.00 %	4.4
Disintegration	DCC-MA-P063/ USP<701>	N.M.T. 15 min	PASSES
Residue on Ignition *	USP	N.M.T. 1.5% Transparent Capsules	PASSES
		N.M.T. 6.0% Colored Capsules	
Identification of HPMC	DCC-MA-P073 / USP	Meets USP Requirements	PASSES
ANALYTICAL			
Arsenic *	EXTERNAL	N.M.T. 0.8 ppm	PASSES
Chromium *	EXTERNAL	N.M.T. 2 ppm	PASSES
Cadmium *	EXTERNAL	N.M.T. 0.5 ppm	PASSES
Lead *	EXTERNAL	N.M.T. 0.5 ppm	PASSES
Mercury *	EXTERNAL	N.M.T. 0.1 ppm	PASSES
Cobalt *	EXTERNAL	N.M.T. 5.0 ppm	PASSES
Vanadium *	EXTERNAL	N.M.T. 10.0 ppm	PASSES
Nickel *	EXTERNAL	N.M.T. 20.0 ppm	PASSES
Total Aerobic Microbial Count	DCC-MA-P031 / USP <61>	N.M.T. 1000 cfu/g	30
Total Yeasts and Molds Count	DCC-MA-P040 / USP <61>	N.M.T. 100 cfu/g	<10
Total Coliforms	DCC-MA-P036 / USP <62>	Absence / 1 g	Absence
Salmonella	DCC-MA-P039 / USP <62>	Absence / 10 g	Absence
Escherichia Coli	DCC-MA-P036 / USP <62>	Absence / 1g	Absence
Staphylococcus aureus	DCC-MA-P037 / USP <62>	Absence / 1g	Absence
Pseudomonas aeruginosa	DCC-MA-P033 / USP <62>	Absence / 1g	Absence

*Reduced Frequency Testing

Storage Conditions: Temperature: 15°C - 30°C / Relative Humidity: 35 % - 70 % RH N.M.T. = No More Than

NOTE: The VISUAL QUALITY is superior to the established figures in the sampling plans of the ANSI/ASQ Z1.4-2013 "Procedure for sampling to inspect for attributes", using simple sampling with level III of General Inspection and Acceptable Level of Quality (AQL) of 0.010 for Critical defects, 0.040 for Major defects and 0.250 for Minor defects. They also fulfill the specifications established in the Technical Information Manual in force.


 Quality Assurance

Date: 2022/11/01



Code: DCC-032G (Valid since November 1st, 2021)
Edition 7

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