

	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	NUVVIT03
Désignation interne	VITAMINE C LIPOSOMALE QUALI C 90 GVT0T

Code client	
Désignation client	VITAMINE C LIPOSOMALE

Numéro de lot	D17654	Numéro de BL	21854
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Date de fabrication	07/01/2024	DDM	01/2027
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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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CONDITIONNEMENT PRODUIT

Type de produit	Gélules HPMC	Couleur	Transparente
PV interne gélules	26152	Taille	Taille 0
Lot Fournisseur gélules	1-001098	Dosage	515mg

Type de conditionnement	Piluliers	Quantité par colis	195*50
Quantité conditionnée	9750	Cartons incomplets	

Autres (dont fond de bol)

COMPOSITION PRODUIT

INGREDIENT(S)	DOSAGE	PV interne	N°LOT FOURNISSEUR
Vitamine C liposomale Quali C	475mg	26123 +26033+26101+26102	06062024-0000000025 + 22052024-00000-00025+17102022-00000-2500E
Amidon de riz	40mg	26132	2420427350

PHYTOCOSMA Laboratoires
23 Route du Burgaud
82600 AUCAMVILLE
Tél. : 05 63 02 71 07
phytocosma@orange.fr
SAS au capital de 13 720,41 euros
RCS MONTAUBAN 432 694 354

06/02/25

Service qualité



CERTIFICATE OF ANALYSIS

Product and Batch Informations			
SUNFLOWER LIPOSOMIAL VITAMIN C			
VITCLIPO_1002			
DV07 -31 05 2024			
Batch	17102022-00000-25000E	Continent of manufacturing	Europe
MF date	17/10/2022	Expiration date	29/10/2025

ANALYSIS ITEM	SPECIFICATION	RESULT	TEST METHOD
Active Ingredients/Substance to control			
Assay	Vitamin C : NLT 67,5%	69,72%	Volumetric determination
Physical/Chemical Control			
Appearance**	White to creamy powder	Complies	Visual inspection
Loss on drying	NMT 5,0%	Complies	105°C - 3 hours
Contaminant Control*			
Lead (Pb)	NMT 3ppm	Complies	Atomic absorption
Arsenic (As)	NMT 1ppm	Complies	Atomic absorption
Cadmium(Cd)	NMT 1ppm	Complies	Atomic absorption
Mercury (Hg)	NMT 0,1ppm	Complies	Atomic absorption
Microbiological Control			
Total aerobic microbial	NMT 20 000 cfu/g	<100cfu/g	As per Eur.Ph
Tot. yeast and mould	NMT 200 cfu/g	<10cfu/g	As per Eur.Ph
Enterobacteria	NMT 100 cfu/g	<10cfu/g	As per Eur.Ph
Salmonella	Negative/25g	Complies	As per Eur.Ph
E.Coli	Negative/g	Complies	As per Eur.Ph
Allegations			
Allergens	Allergen FREE		
GMO	No GMOs		
Irradiation	No irradiation		
BSE/TSE	BSE/TSE FREE		
Nanoparticles	Nanoparticle FREE		
Vegans/vegetarians	Suitable		
Packing and Storage			
Packing	All requirements for Food Regulation		
Storage	Store in its closed original packaging at room temperature (5-25°C), away from heat, direct light and moisture		

*According to a control plan

**Depending to variations of raw material



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Tel : +33 241 475 225 Fax : +33 241 476 044
Mail : quality@natural-ingredients.fr

29/10/2024
Angebault Mélanie
Quality Department

CERTIFICATE OF ANALYSIS

Product and Batch Informations

LIPOSOMAL VITAMIN C SUNFLOWER

REF : VITCLIPO_1002

DT V07 - 31/05/2024

Batch	22052024-00000-00025	Origin (natural/synthetic)	Natural
N° CAS	50-81-7	Country of manufacturing	Europe
MF date	22/05/2024	Expiration date	21/05/2026

ANALYSIS ITEM	SPECIFICATION	RESULT	TEST METHOD
Active Ingredients/Substance to control			

Assay	Vitamin C content : NLT 67.5%	68,10%	LC DAD
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Physical/Chemical Control			
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Appearance	White to creamy powder	Complies	Visual inspection
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Loss on drying	NMT 5%	Complies	105°C - 3 hours
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Contaminant Control*			
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Lead (Pb)	NMT 3ppm	Complies	Atomic Absorption
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Arsenic (As)	NMT 2ppm	Complies	Atomic Absorption
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Cadmium(Cd)	NMT 1ppm	Complies	Atomic Absorption
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Mercury (Hg)	NMT 0,1ppm	Complies	Atomic Absorption
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Microbiological Control			
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Total aerobic microbial	NMT 20 000 cfu/g	Complies	As per Eur.Ph
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Tot. yeast and mould	NMT 200 cfu/g	Complies	As per Eur.Ph
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Salmonella*	Negative/25g	Complies	As per Eur.Ph
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E.Coli*	Negative/g	Complies	As per Eur.Ph
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Statements			
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Allergens	Allergen free
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GMO	No OGM
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Irradiation	No irradiation
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BSE/TSE	BSE/TSE Free
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Nanoparticles	Nanoparticles Free
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Vegans/Vegetarians	Suitable for vegans/vegetarians
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Packing and Storage			
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Packing	Suitable for food industry
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Storage	Store in dry places and keep away from strong direct light and heat.
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*According to a control plan

CERTIFICATE OF ANALYSIS



Product and Batch Informations

LIPOSOMAL VITAMIN C SUNFLOWER

REF : VITCLIPO_1002

DT V07 31/05/2024

Batch	06062024-00000-00025		
N° CAS	50-81-7	Country of manufacturing	Europe
MF date	06/06/2024	Expiration date	05/06/2026

ANALYSIS ITEM	SPECIFICATION	RESULT	TEST METHOD
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Active Ingredients/Substance to control

Assay	NLT 67,5% Vitamin C content	71,40%	LC DAD
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Physical/Chemical Control

Appearance	White to creamy powder	Complies	Visual inspection
Loss on drying	NMT 5%	Complies	105°C - 3 hours

Contaminant Control*

Lead (Pb)	NMT 3ppm	Complies	Atomic Absorption
Arsenic (As)	NMT 2ppm	Complies	Atomic Absorption
Cadmium(Cd)	NMT 1ppm	Complies	Atomic Absorption
Mercury (Hg)	NMT 0,1ppm	Complies	Atomic Absorption

Microbiological Control

Total aerobic microbial	NMT 20 000 cfu/g	Complies	As per Eur.Ph
Tot. yeast and mould	NMT 200 cfu/g	Complies	As per Eur.Ph
Salmonella*	Negative/25g	Complies	As per Eur.Ph
E.Coli*	Negative/g	Complies	As per Eur.Ph

Statements

Allergens	Allergen free
GMO	No OGM
Irradiation	No irradiation
BSE/TSE	BSE/TSE free
Nanomaterials	Nanomaterials free
Vegans/ Vegetarians	Suitable for vegans and vegetarians

Packing and Storage

Packing	Suitable for food industry
Storage	Store in dry places and keep away from strong direct light and heat.

*According to a control plan



NATURAL

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14/11/2024

Maeva FOURNY

Quality Department

General Information

Product	Remy O DR6	Production Date	31/07/2024 (dd/mm/yyyy)
Batch	2420427350	Best before	30/07/2028 (dd/mm/yyyy)
Issued by	Quality Control Management	Date CoA Issued	26/09/2024 (dd/mm/yyyy)

Results of analyses

Parameter	Result	Unit	Method ⁽¹⁾	LSL	USL
Physical and Chemical Parameters					
Moisture	6	g/100g	ISO 712	≤	14
Protein content (N*6,25) on DM	4,4	g/100g d.m.	ISO 1871 ⁽¹⁾	≤	7,0
Ash content on DM	0,2	g/100g d.m.	ISO 3593	≤	1,0
Rheological Parameters					
Starting gel point, pH as is, 6%	82	°C	Brabender	≥	60
End viscosity, pH as is, 6%	640	BU	Brabender	≥	500
Microbiological Parameters					
Salmonella (/375g)	Negative	/375g	ISO 6579		Negative
Total mesophilic bacteria (aerobic)	20.000	cfu/g	ISO 4833	≤	100.000
Yeasts and Moulds	270	cfu/g		≤	1.000
Enterobacteriaceae	80	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

CERTIFICATE OF ANALYSIS

EMPTY HARD CAPSULES OF VEGETABLE ORIGIN

CUSTOMER: GOCAPS GMBH			
LOT No.:	K2408001649	PRODUCT CODE:	K00003G
PURCHASE ORDER NUMBER:	PO2000837	CHARGE No.:	1-001098
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	%
<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. 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>			

Date of Manufacture: 2024-09

Expiration Date: 2029-09

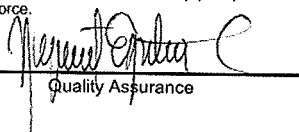
CRITERIA	METHOD / REFERENCE	SPECIFICATIONS	RESULTS
PHYSICAL			
Average Capsule Weight	DCC-MI-P003/ USP<2091>	103.00-115.00 mg	105.4
Loss on drying	DCC-MA-P027	3.00-8.00 %	4.7
Disintegration	DCC-MA-P063/ USP<701>	N.M.T. 15 min	PASSES
Residue on Ignition *	USP	N.M.T. 1.5% Transparent Capsules	PASSES
		N.M.T. 6.0% Colored Capsules	PASSES
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	10
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 of 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.

Approval by:


Quality Assurance

Date: 2024/10/09



Code: DCC-032G (Valid since March 15th 2024)
Edition 9

MANUFACTURER ADDRESS: VIA 40 85-48 BARRANQUILLA - COLOMBIA
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