	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	D17088	Numéro de BL	21061+21065
Date de fabrication	30/01/2024	DDM	01/2026

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

Le produit contient de(s) Allergène(s) : Non

Liste des allergène(s) dans le produit

Non applicable

Le produit contient de(s) Additif(s) : Non

Liste de(s) additif(s) dans le produit

Non applicable

Le produit est BIO : Non

*(*Produit issu de l'algriculture biologique FR-BIO-01 et process conforme à la fabrication de produits biologiques)*

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

CONDITIONNEMENT PRODUIT

Type de produit	Gélules HPMC	Couleur	Transparente
PV interne gélules	25160	Taille	Taille 0
Lot Fournisseur gélules	1-000628	Dosage	515mg

Type de conditionnement	Piluliers	Quantité par colis	7*119+9*119
Quantité conditionnée	6852	Cartons incomplets	96*50 +1*67+1*81

Autres (dont fond de bol)	

COMPOSITION PRODUIT

INGREDIENT(S)	DOSAGE	PV interne	N°LOT FOURNISSEUR
Vitamine C liposomale Quali C	475mg	25195	15092023-00000-00025
Amidon de riz	40mg	24992+25123	23202960020+2320326080

08/04/24
Service qualité

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

CERTIFICATE OF ANALYSIS

Product and Batch Informations			
VITAMINE C LIPOSOMAL REF : VITCLIPO_1002 DT V05 - 21/12/2021			
Batch	15092023-00000-00025	**Origin (natural/synthetic)**	Synthetic
N° CAS	50-81-7 (Ascorbic acid)	**Country of manufacturing**	Europe
MF date	15/09/2023	**Expiration date**	14/09/2025

ANALYSIS ITEM	SPECIFICATION	RESULT	TEST METHOD
Active Ingredients/Substance to control			
Assay	Vitamin C : NLT 67,5%	Complies	Volumetric determination
Physical/Chemical Control			
Appearance**	White to creamy powder	Complies	Visual inspection
Sieve analysis	NLT 95,0% through 40 mesh	Complies	Circular Vibrating Screener
	NLT 70,0% through 60 mesh	Complies	
Loss on drying	NMT 5,0%	Complies	105°C - 3 hours
Contaminant Control*			
Heavy metals	NMT 10ppm	Complies	Atomic absorption
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	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*

***Sum of ethylene oxide and 2-chloro-ethanol expressed as ethylene oxide*

****According to meeting on Ethylene oxide on october 2021, applicable limit of ethylene oxide for food supplement, some minerals and additives (except in specific case) is 0.1ppm*

NATURAL
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Tel :+33 241 475 225 Fax :+33 241 476 044
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19/10/2023
GANDILHON Damien
Quality Department

connecting nutrition and health

Certificate of Analysis

General Information

Product	Remy O DR6	**Production Date**	16/05/2023	(dd/mm/yyyy)
Batch	2320326080	**Best before**	15/05/2027	(dd/mm/yyyy)
Issued by	Quality Control Management	**Date CoA Issued**	06/06/2023	(dd/mm/yyyy)

Results of analyses

Parameter	Result	Unit	Method [1]	LSL	USL
Physical and Chemical Parameters					
Moisture	12	g/100g	ISO 712		≤ 14
Protein content (N*6,25) on DM	5,1	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%	81	°C	Brabender	≥ 60	
End viscosity, pH as is, 6%	611	BU	Brabender	≥ 500	
Microbiological Parameters					
Salmonella (/375g)	Negative	/375g	ISO 6579		Negative
Total mesophilic bacteria (aerobic)	34 000	cfu/g	ISO 4833		≤ 100 000
Yeasts and Moulds	70	cfu/g			≤ 1 000
Enterobacteriaceae	90	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

CapsCanada®
a Lyfe Group Co.

K-CAPS®
Capsules of Vegetable Origin

CERTIFICATE OF ANALYSIS
EMPTY HARD CAPSULES OF VEGETABLE ORIGIN

CUSTOMER: GOCAPS GMBH
LOT No.: K2307002420 **PRODUCT CODE:** K00003 **SIZE:** 0
PURCHASE ORDER NUMBER: PO2000470 **CHARGE No.:** 1-000628 **ART No.:** 56-000107
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	%

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.

Date of Manufacture: 2023-08 Expiration Date: 2028-08

CRITERIA	METHOD / REFERENCE	SPECIFICATIONS	RESULTS
PHYSICAL			
Average Capsule Weight	DCC-MI-P003/ USP<2091>	103.00-115.00 mg	107.9
Loss on drying	DCC-MA-P027	4.00-8.00 %	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 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	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 Frecuency 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.

Quality Assurance

Date: 2023/09/08

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

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