	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	NUVVIT05
Désignation interne	VITAMINE C QUALI
Code client	
Désignation client	VITAMINE C

Numéro de lot	D17655	Numéro de BL	21568
Date de fabrication	11/12/2024	DDM	12/2026

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	26166	Taille	Taille 1
Lot Fournisseur gélules	1-001090	Dosage	505mg

Type de conditionnement	Piluliers	Quantité par colis	120*50
Quantité conditionnée	6004	Cartons incomplets	1*4

Autres (dont fond de bol)	

COMPOSITION PRODUIT

INGREDIENT(S)	DOSAGE	PV interne	N°LOT FOURNISSEUR
Vitamine C L-ascorbique Quali C	500mg	26117	1124054014
Amidon de riz	5mg	26008	2420427350

18/12/24
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

06
1.10.24
A.Wait

COVITAS 可维素
HEILONGJIANG NHU BIOTECHNOLOGY CO., LTD.
No.2 Haotian Road, Economic and Technological Development Zone, Suihua, Heilongjiang, China
Tel:(+86)575-86133360 Fax:(+86)575-86165366 Http: // www.cnhu.com

Certificate of Analysis

Contract No.: 803947 Report No.: 20240127
Invoice No.: 300878 Container No.:OOCU9166441

Product Name	Ascorbic Acid	Product Code	SPVC06
Batch No.	1124054014	Manufacture Date	2024-05-16
Quantity	1200kg	Analysis Date	2024-05-18
Foundation	USP42/EP11.0/EU231/2012	Expiry Date	2027-05-15

Results of Analysis:

Items	Standards	Results
Appearance	White or almost white, crystalline powder or colourless crystals	Qualified
Identification	Positive Reaction	Qualified
Assay（$C_6H_8O_6$）	99.0% ~ 100.5%	99.9%
Specific optical rotation	+20.5°~+21.5°	+20.81°
pH(with 2% water solution)	2.4~2.8	2.6
pH(with 5% water solution)	2.1~2.6	2.4
Clarity of solution	Clear	Clear
Colour of solution	≤BY_7	<BY_7
Melting point	190~192℃	190.3-191.4℃
Particle size	≥95% through 100mesh	98.8%
Residue on ignition**	≤0.1%	<0.1%
Heavy metal**	≤10.0 mg/kg	Conform
Iron**	≤2.0 mg/kg	Conform
Copper**	≤5.0 mg/kg	Conform
Lead**	≤0.5 mg/kg	Conform
Arsenic**	≤1.0 mg/kg	Conform
Mercury**	≤0.1 mg/kg	Conform
Cadmium(Cd)**	≤0.5 mg/kg	Conform
Impurity E*	≤0.20%	Conform
Impurities C，D*	≤0.15%	Conform
Unspecified impurities (each impurity)*	≤0.10%	Conform
Sum of impurities other than C and D*	≤0.20%	Conform
Residual Solvents (as methanol)*	≤200 mg/kg	Conform
Loss of Drying*	≤0.4%	Conform
Total Plate Counts*	≤1000 cfu/g	Conform
Yeasts and Moulds*	≤100 cfu/g	Conform
Escherichia coli*	Negative in 10g	Conform
Salmonella*	Negative in 25g	Conform
Staphylococcus Aureus*	Negative in 25g	Conform

* Test once every three months.

** Test once every six months..

Reporter: 李春雨 Reviewer: 王伟杰

Approver: 郑乐友

QC Manager: Leyou Zheng

HEILONGJIANG NHU BIOTECHNOLOGY CO., LTD. QC

connecting nutrition and health

Certificate of Analysis

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

This certificate is issued automatically, it does not need a signature .

FORM Beneo-Remy QC lot Coa V:02

LIMS: 117.731

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.: K2408001642 **PRODUCT CODE:** K00016G **SIZE:** 1

PURCHASE ORDER NUMBER: PO2000829 **CHARGE No.:** 1-001090 **ART No.:** 56-000992

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: 2024-09 Expiration Date: 2029-09

CRITERIA	METHOD / REFERENCE	SPECIFICATIONS	RESULTS
PHYSICAL			
Average Capsule Weight	DCC-MI-P003/ USP<2091>	75.00-85.00 mg	78.3
Loss on drying	DCC-MA-P027	3.00-8.00 %	4.3
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	9
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.

Approval by: _____
Quality Assurance

Date: 2024/09/25

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

MANUFACTURER ADDRESS: VIA 40 85-48 BARRANQUILLA - COLOMBIA
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