THE RAPTOR COMPANY

65, rue Réaumur 75002 PARIS

<u>A l'attention de</u> Monsieur Ismail OUSLIMANI

Paris, le 9 octobre 2023

Objet: ELEMENTS

LIBERATION DES LOTS Service Qualité Certification Monsieur,

Nous vous confirmons la libération du lot de ELEMENTS numéro F-23-111 en date du 06/07/2023.

Ce lot est certifié conforme en tous points, et est libéré pour sa commercialisation.

Nous attestons que l'ensemble de ce lot a été produit en France et vous remercions de bien vouloir trouver en pièce(s) jointe(s) :

MASTER FILE PRODUIT

Nous vous remercions pour votre confiance, Bien cordialement,

Le Pharmacien Responsable Qualité

THE RAPTOR COMPANY RAPTOR NUTRITION

ELEMENTS

PRODUCT MASTER FILE

Direction générale de l'alimentation

Liberté Égalité Fraternité

DIRECTION GÉNÉRALE DE L'ALIMENTATION 251 RUE DE VAUGIRARD 75732 PARIS CEDEX 15

BEPIAS (Bureau des Etablissements et Produits des Industries Alimentaires Spécialisées)

Tél : 01 49 55 84 17 / 01 49 55 84 02 Mél : <u>bepias.sdssa.dgal@agriculture.gouv.fr</u> Paris, le 20/07/2023

ATTESTATION DE DECLARATION D'UN COMPLEMENT ALIMENTAIRE

La Direction générale de l'alimentation (DGAL) atteste que :

THE RAPTOR COMPANY

a effectué, le 28/06/2023, la déclaration mentionnée à l'article 15 du décret n°2006-352 du 20 mars 2006 relatif aux compléments alimentaires pour le produit :

ELEMENTS multivitamines & mineraux Gélule

Cette déclaration est enregistrée sous le numéro : 2023-6-1430

La déclaration prévue à l'article 15 du décret n°2006-352 vise à informer l'administration de la mise sur le marché d'un complément alimentaire.

Votre produit est enregistré sur la liste consultable à l'adresse suivante : https://teleicare.dgccrf.finances.gouv.fr/ Home/ConsulterAttestation. Il pourra faire l'objet de contrôles de l'administration.

Cette attestation ne constitue donc pas une garantie de conformité aux dispositions en vigueur ni une autorisation de mise sur le marché..

Conformément aux dispositions de l'article 17 du règlement (CE) n° 178/2002 établissant les principes généraux et les prescriptions générales de la législation alimentaire, il vous incombe de veiller à ce que le complément alimentaire que vous mettez sur le marché réponde à la définition des compléments alimentaires au sens de l'article 2 du décret n°2006-352, ainsi qu'aux prescriptions du droit alimentaire qui leur sont applicables, notamment en matière d'hygiène (R(CE) n°852/2004), d'information du consommateur (R(UE) n°1169/2011) et de nouveaux aliments (R(UE) n°2015/2283). À cet égard, vous devez être en mesure de prouver que les ingrédients mis en œuvre disposent d'un historique de consommation dans l'Union européenne avant le 15 mai 1997 ou, à défaut, qu'ils ont été autorisés au titre du règlement (UE) n°2015/2283.

Pour tout renseignement d'ordre général sur le cadre réglementaire applicable aux compléments alimentaires, rendez-vous sur le site de la DGAL à l'adresse suivante : https://agriculture.gouv.fr/quest-ce-quun-complement-alimentaire.



Flasher le QR-Code ou saisir le code de vérification du document **MZnbiZ** sur le site <u>MZnbiZ</u>

Dossier de lot F-23-111 ELEMENTS / RAPTOR NUTRITION

Général	
Date début de production :	05/07/2023
Date fin de production :	06/07/2023
DLUO:	19/06/2026

Préparation	
Quantité demandée :	125 000
Quantité total engagée :	125 000
Perte :	0 %

Contenant		
Qualité :	НРМС	
Taille :	T.0	
Couleur :	TR.	

Composition

Ingrédient	Quantité / Unité	Perte	Quantité à produire	Quantité prélevée	N° Lot
CH-Calcium Citrate (24%)	120 Mg	0 %	15 000 g	15 000 g	312207111
CH-Potassium citrate (36%)	350 Mg	0 %	43 750 g	43 750 g	322210211
CH-Fer bisglycinate 20%- 23%	32,91 Mg	0 %	4 113,75 g	4 113,75 g	DHJLYT2209061
VIT-Vit. E-Nat 700 UI (45%)	13,5 Mg	0 %	1 687,5 g	1 687,5 g	A(3941020-1249)EG/7
VIT-Vit. PP (ou B3- NiacinNicotinam.)	9,6 Mg	0 %	1 200 g	1 200 g	b2208nia156r
VIT-Vit. A (Rétinol acétate 500.000 UI/g) 12-14%	3,325 Mg	0 %	415,63 g	415,63 g	0-2203032-210417
VIT-Vit. B8 (ou H- Biotine)	0,03 Mg	0 %	3,75 g	3,75 g	CS1-2110022
CH-Manganese gluconate (11.4%)	8,772 Mg	0 %	1 096,5 g	1 096,5 g	20210226210
CH-lodure potassium (77% lode 23.55% de K)	0,097 Mg	0 %	12,18 g	12,18 g	112320917
CH-Chrome Picolinate (12,2%)	0,163 Mg	0 %	20,31 g	20,31 g	16102021
CH-Cuivre gluconate (14%)	3,572 Mg	0 %	446,44 g	446,44 g	20210523
CH-Sélénométhionine (0.5%)	5,5 Mg	0 %	687,5 g	687,5 g	O-1602163-220105
VIT-Vitamine K2-MK7 (1%)	7,5 Mg	0 %	937,5 g	937,5 g	2204M05
AD- BIO Amidon de riz	75 Mg	0 %	9 375 g	9 375 g	221006
Total (Mise en Oeuvre)	629,97 Mg		78 746,05 g		

Contenant	Quantité	N Lot Fourn.
G-Vg-0-Transp	125 000	SV022081330

Fabrication

Informations	
Opérateur :	ВЕ
Vitesse :	1800

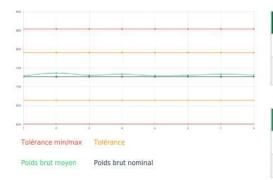
Poids		
Poids net :	630 Mg	
Contenant poids :	98 Mg	
Poids brut nominal :	728 Mg	

Tolérance		
Tolérance / Poids net :	-10% / +10%	
Soit poids unitaire brut :	>665Mg & <791Mg	

Relevés

Contrôle Poids Bruts moyens (Lot de 10 gélules): 730,737,731,734,730,731,734,732

Dossier de lot F-23-111 ELEMENTS / RAPTOR NUTRITION



Poids moyen sur 10 gélules (Mesure 8)		
Poids brut : 732 Mg		
Poids net :	634 Mg	

Contrôle Uniformité de masse : Mesures : 80			
< 665 Mg : (0 %)	< 791 Mg : (0 %)		
< 602 Mg : (0 %)	< 854 Mg : (0 %)		

Quantité théorique à obtenir :	124 205
Quantité unité reg. machine :	
Quantité nette à obtenir :	124 205

Quantité produite :	122 400
Rendement :	99%

Tricalcium citrate

Product SPECIFICATION

Product Name Tricalcium citrate CAS Code 5785-44-4

The product meets OR exceeds: USP32	
Appearance:	white, powder, crystalline
Identification	A to pass test, B to pass test
Assay %	97.5-100.5%
Fluoride, ≤mg/kg	30
Heavy Metal ≤%	0.002
Lead, ≤mg/kg	3
Mercury, ≤mg/kg	0.1
Loss on drying %	10.0-13.3
Acid insoluble substances, ≤%	0.2
Arsenic (as As), ≤mg/kg	1
Cadmium (Cd), ≤mg/kg	1
Organic volatile impurities	Meets requirements

Tricalcium citrate

BSE/TSE Statement

The product does not contain and is not derived from Specified Risk Material as defined in Commission Directive European and it is conform to the EU legislation relating to the risk of Transmission of Spongiform Encephalopathy (TSE).

GMO Statement

Following EU regulations have been published in the Official Journal of the European Union:

- Regulation (EC) No.1829/2003 of the European Parliament and the Council of 22nd September 2003 on genetically modified food and feed
- Regulation (EC) No.1830/2003 of the European Parliament and the Council of 22nd September 2003 concerning tractability and labeling on genetically modified organisms and tractability of food and feed products produced from genetically modified organisms and amending Directive 2001/18/EC.

This product has not been genetically modified so there is no obligation of GMO labeling as defined by the above mentioned regulations.

NON IRRADIATION/NON IONIZED Statement

According to directives 1992/2/EC and 1999/3/EC, We confirm is not made from irradiated/ionized raw materials or was irradiated/ionized the product.

Tricalcium citrate

PESTICIDES Statement

To our actual knowledge of production process, raw materials and equipment used potential pesticide residues in the above mentioned product comply with the European legislation on pesticide residues, esp. Regulation (EC)None. 396/2005. However, we do no test for pesticide residues on a routine basis.

RESIDUAL SOLVENTS Statement

The product meets with:

- DIRECTIVE 2009/32/EC OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 23 April 2009 on the approximation of the laws of the Member States on extraction solvents used in the production of foodstuffs and food ingredients
- DIRECTIVE 2010/59/EU of 26 August 2010 amending Directive 2009/32/EC of the European Parliament and of the Council on the approximation of the laws of the Member States on extraction solvents used in the production of foodstuffs and food ingredients.

CONTAMINANTS

The product is in accordance and meets with the commission regulation (EC) No.629/2008 of 2 July 2008 amending regulation (EC) No.1881/2006 setting maximum levels for certain contaminants in foodstuffs.

PCBS and MELAMINE

The product is in accordance and meets with the commission regulation (EU) No.594/2012 of 5 July 2012 amending regulation (EC) No.1881/2006 as regards the maximum levels of the contaminants Ochratoxin a, non dioxin-like PCBS and melamine in foodstuffs.

ANTI-DOPING

As per the list of 2012 of the world anti-doping agency the ingredient is not a doping substance and not a combination of doping substances. The ingredient does not contain any doping substance. The ingredient does not result from a doping substance.

NANOMATERIALS Statement

According to the the definition of 'nanomaterials' of the UE REGULATION (EU) No 1169/2011 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 25 October 2011 on the provision of food information to consumers , we hereby attest that no nanomaterials are used in the formulation nor in the packaging materials of our product.

Tricalcium citrate

ALLERGENS Statement

In accordance with the Directive 2003/89/EC and Council amending Directive 2000/13/EC and taking Directive 2005/26/EC, Directive 2005/63/EC and Directive 2006/142/EC of 22.12.2006 into consideration. We confirm the following information:

Note: Does the raw material CONTAIN or is it DERIVED from or could it be CROSS-CONTAMINATED by any of the following.

Cereals containing gluten (i.e. wheat, rye, barley, oats, spelt, kamut or their hybridised strains) and products thereof	No
Crustaceans and products thereof	No
Eggs and products thereof	No
Fish and products thereof	No
Peanuts and products thereof	No
Soybeans and products thereof	No
Milk and products thereof (including lactose)	No
Nuts (i.e. Almond, Hazelnut, Walnut, Cashew, Pecan nut, Brazil nut, Pistachio nut, Macadamia nut and Queensland nut and products thereof)	No
Celery and products thereof	No
Mustard and products thereof	No
Sesame seeds and products thereof	No
Sulphur dioxide and sulphite at concentrations of more than 10 mg/kg or 10 mg/l expressed as SO2	No
Lupine and products thereof	No
Molluscs and products thereof	No

Potassium citrate tribasic

Product SPECIFICATION

Product Name	Potassium citrate tribasic
CAS Code	6100-05-6

Description	White, hygroscopic, granular powder or transparent crystals	
Identification	Passes test	
Loss on drying, %	≤6.0	
Assay, %	≥99.0	
Oxalates, %	≤0.01	
Heavy metal (As Pb), ≤mg/kg	≤5	
рН	7.5-9.0	
Arsenic (As), ≤mg/kg	≤1	
Lead (Pb), ≤mg/kg	≤3	
Mercury (Hg), ≤mg/kg	≤0.1	
Cadmium (Cd), ≤mg/kg	≤1	

Potassium citrate tribasic

BSE/TSE Statement

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GMO Statement

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Potassium citrate tribasic

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Eggs and products thereof	No
Fish and products thereof	No
Peanuts and products thereof	No
Soybeans and products thereof	No
Milk and products thereof (including lactose)	No
Nuts (i.e. Almond, Hazelnut, Walnut, Cashew, Pecan nut, Brazil nut, Pistachio nut, Macadamia nut and Queensland nut and products thereof)	No
Celery and products thereof	No
Mustard and products thereof	No
Sesame seeds and products thereof	No
Sulphur dioxide and sulphite at concentrations of more than 10 mg/kg or 10 mg/l expressed as SO2	No
Lupine and products thereof	No
Molluscs and products thereof	No

Ferrous Glycinate

	SPECIFICATION D	ATA		
Nom chimique / Chemical Name	Ferrous bisglycinate			
Numéro CAS / CAS number	20150-34-9	20150-34-9		
Numéro EINECS / EINECS number	606-444-7			A111
Masse Molaire / Molecular mass	239.99 g/mol	239.99 g/mol		
Formule brute / Molecular Formula	Fe(O ₂ C ₂ H ₂ NH ₂) ₂ .2H ₂ O			H ₂ N 0. Fe ⁺⁺
Conformité / Compliance	Internal standard			. 0 0 0
Utilisation possible / Use	Alimentaire,	Food, food		
	complément	supplement		
	alimentaire			
Apparence / Appearance	Poudre		Powder	
Odeur, couleur, goût / Odor, color, taste	Marron foncé ou gris-ver	rt	ART ARTER SOUSH	wn or grey-green
Granulométrie / Particle size	≥ 80 mesh		Dark brow	wit of grey-green
Solubilité / Solubility	≥ 00 IIIesii			
Densité / Bulk density	0.65 g/ml			
Excipient / Carrier	Aucun		None	
Excipient / Carrier	Méthode d'analyses	Fufavones		S- f-ifi-ations
	Analysis method	Fréquence of Frequency of	f analysis	Spécifications Specifications
Teneur/ Assay (on dried basis)	GB – titration analysis	Chaque lot / E		(Fe ²⁺): 20.0 – 23.7%
Perte à la dessication/ Loss on drying	GB	Chaque lot / E	ach batch	≤7.0%
Cendres sulfuriques /Sulphated ashes				
рН				
Pouvoir rotatoire / Specific rotation				
Point de fusion / Melting point				
Identification / Identification				
Nitrogen content (on dried basis)	GB	Chaque lot / E	ach batch	10.0 – 12.0%
Ferric ion (Fe ³⁺) (on dried basis)	GB – titration analysis	Chaque lot / E	ach batch	≤2.0%
Total iron	GB – titration analysis	Chaque lot / E	ach batch	19.0 – 24.0%
Métaux lourds totaux / Total heavy metals	1	6		
Arsenic / Arsenic	GB	Annuellement	/ Yearly	≤1 ppm
Plomb / Lead	GB	Chaque lot / E		≤1 ppm
Cadmium / Cadmium	GB	Annuellement		≤1 ppm
Mercure / Mercury	GB	Annuellement	SECURIO CONTRACTOR	≤ 0.1 ppm
Aluminium / Aluminium			The second	P.P.
Auditinativ Phalimian		k		
Flore totale / Total plate count (CFU/g)	GB	Annuellement	/ Yearly	≤ 1 000
Levures et moisissures/Yeast & mold (CFU/g)	GB	Annuellement	/ Yearly	≤ 100
Enterobacteria				
E. coli	GB	Annuellement	/ Yearly	Negative
Salmonella	GB	Annuellement	/ Yearly	Negative
Staphylococcus aureus	GB	Annuellement	/ Yearly	Negative
Listeria monocytogenes			7.7	_
Is the product a part of the monitoring plan?	No Yes for som	ne microbiologica	and heavy	metals parameters.

Ferrous Glycinate

Conditions d'entreposage/Storage conditions (T°)	Stocker dans un emballage fermé, à température ambiante < 25°C à l'abri de la lumière, de l'humidité et de sources de chaleur. / Keep in well-closed containers under room temperature < 25°C away from light, humidity and heat sources.
Conditionnement / Packaging	Sterile PE bag inside, outsourced to carton or paper bucket
Durée de conservation / Shelf life	2 ans / 2 years

GENETICALLY MODIFIED ORGANISMS - GMO

According to EU regulations EC/1829/2003 & EC/1830/2003

According to the EU regulations concerning the traceability and labelling of Genetically Modified Organisms (GMO), and the traceability of food products produced from GMO (Regulation N°1829/2003/EC and N°1830/2003/EC and amendments), we confirm that the product:

produc	ι.			
1.	>	Contains ingredient(s) from Maize and/or Soya origin	Yes 🗌	No 🖾
	Contains no material which contains, consists of or is produced from authorized GMOs in proportion higher than 0,9 %. This presence is adventitious or technically unavoidable		Yes 🖂	No 🗆
	>	is GMO or contains or consists of GMOs	Yes 🗌	No 🖾
	>	is produced from GMOs or contains ingredients produced from GMOs	Yes 🗌	No 🖾
2.	>	Contains ingredient(s) from other potential GMO-risk raw material(s) (rice, beet, cotton)	Yes 🔲	No 🖾
	If yes:	 ingredient: vegetable origin: geographic origin: Contains no material which contains, consists of or is produced from evaluated		
		GMOs in proportion higher than 0,5 % . This presence is adventitious or technically unavoidable	Yes 🔼	No 🗆
3.	with GM derived f	mation about processing aids, is produced or contains ingredient(s) produced Os or Genetically Modified Microorganisms processing aids (bacteria, yeast) or rom GMOs or GMMs (including enzymes)	Yes 🗌	No 🖾
	If yes, ple	ease specify:		

	F-3-44-4-4-7	St		
	rerrous C	Slycinate		
	NANOM	IFTADIAI		
NANOMETARIAL According to regulation EU N°1169/2011				
Does this raw material contain or can be considered as an engineered nanomaterial*? Was this raw material made with nanotechnology? Does the product contain any nanotechnology as defined by the French decree n°2012-232? Information based on: Analysis Supplier declaration Other			YES	NO⊠ NO⊠ NO⊠
or that is composed of discrete f	unctional parts, either internally o g structures, agglomerates or aggr	erial that has one or more dimensi r at the surface, many of which ha regates, which may have a size abo	ve one or more dimensi	ons of the
	RESIDUAL	SOLVENTS		
According to	UE Directive 2009/32 modifie	d and 20010/59 and MGP/ICH	/283/95 Q3C	
Does the raw material contain If yes: - percentage and name	n any residual solvent? e of each solvent used:		YES	NO⊠
Is the product in compliance with the French decree 27 August 2009? YES NO VES NO				NO NO NO
)	FOOD CE	RTIFICATE		
Is the product complying with commission regulation (EC) N° 178/2002 laying down the general principles are requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety and with the regulations 2073/2005 and 852/2004 on food hygiene? Manufacturer's quality certificates: ISO 9001 SISO 14001 FSSC 22000 ISO 22000 HACCP BRC IFS FAMI QS GMP Other: ISO 45001 FDA registration number:				atters o
ССР	Location on process	Limit	Corrective acti	on
CCP1	Raw material receiving	According to the related SOP	Refused to accept	
Was the product used in the European food industry before May 1997 following the regulations 258/97 and 2015/2283? YES NO If the product is an authorized Novel Food, does it comply with the requested specifications from the regulation 2015/2283? YES NO For food supplement only			NO NO	
is the raw material rood drade:				

D-ALPHA TOCOPHERYL ACETATE (700 UI-G) (VIT E NAT)

PRODUCT SPECIFICATION					
Product Name	INCI Name	EINECS Number	CAS Code		
D-ALPHA TOCOPHERYL ACETATE (700 UI-G) (VIT E NAT)	alpha-tocopheryl acetate	200-405-4	58-95-7		
Names			emical name: d-alpha tocopheryl acetate , 4´R, 8´R-alpha-tocopheryl acetate		
INCI name		toc	opheryl acetate		
General Characterization					
Off-white/beige free flowing gra	anular powder.				
	E acetate is derived from non-C y physical and chemical means				
It is then absorbed onto silicon the food and dietary suppleme	dioxide. It is intended for use a nt industry.	s Vitamin E in			
Identification					
Name		Em	pirical formula		
d-alpha tocopheryl acetate		C3	1H52O3		
Ingredients/Composition		GC	method according to IT-02-L		
D-alpha Tocopheryl acetate*		514	1.7 mg/g min.		
Vegetal (sunflower) oil:		95	mg/g min		
Silicon dioxide SiO2 carrier (fr	ee flowing type)	390	0.3 mg/g max		
*Derived from soybean oil					
Quality Control Assay		(Pa	rameters analyzed per batch)		
Appearance		Off	-white to beige granular powder		
Taste		Bla	nd, characteristic		
Odour		Bla	nd to none		
Specific gravity/Density (25°C)		0.5	0 – 0.80 g/cm3		
Potency		700	700 I.U/g (min)		

D-ALPHA TOCOPHERYL ACETATE (700 UI-G) (VIT E NAT)

PRODUCT SPECIFICATION	
Acidity (as d-α-tocopheryl acetate)	< 0.35 ml KOH 0.1N /g
Loss on drying	< 5 %
Benzo (a) pyrene	< 2 ppb
Sum of Benzo(a)pyrene, Benzo(a)anthracene, Benzo(b)fluoranthene and Chrysene	< 10 ppb
(Parameters analyzed periodically based on statistical data	
Particle size	90 % ≥ 400 µm (40 mesh)
Heavy metals (for D-alpha-tocopheryl acetate)	Pb: <0.1ppm Hg:<0.1ppm Cd: <1ppm As: <1ppm
Heavy metals (for Silicon dioxide):	Pb: <5ppm Hg:<1ppm Cd: <2ppm As: <3ppm
Microbiological tests	(In compliance with EP and RD 3177/83)
E. Coli	None cfu/g
Salmonella	None cfu/25g
Lysteria Monocytogenes	None cfu/10 g
Shigella	None cfu/10 g
Clostridium Perfringens	None cfu/g
Staphylococcus Aureus	None cfu/g
Enterobacteriaceae	max 100 cfu/g
Total Aerobic Count	max 1000 cfu/g
Yeast & Moulds	max 100 cfu/g
Residual solvents	In compliance with limits established in EP and USP for Class I solvents. This product meets as well Directive 2009/32/EC

D-ALPHA TOCOPHERYL ACETATE (700 UI-G) (VIT E NAT)

BSE/TSE

This product does not contain and is not derived from Specified Risk Material as defined in European Commission Directive (and its amendments) and is therefore conform with the EU legislation relating to the risk of Transmission of Spongiform Encephalopathy (TSE).

GMO

The following EU regulations have been published in the Official Journal of the European Union in this subject:

- Regulation (EC) No.1829/2003 of the European Parliament and the Council of 22 September 2003 on genetically modified food and feed.
- Regulation (EC) No.1830/2003 of the European Parliament and the Council of 22 September 2003 concerning tractability and labeling on genetically modified organisms and tractability of food and feed products produced from genetically modified organisms and amending Directive 2001/18/EC.

This product has not been genetically modified so there is no obligation of GMO labeling as defined by the above mentioned regulations.

NON IRRADIATION/NON IONIZED

According to directives 1992/2/EC and 1999/3/EC, we confirm that this product is not made from irradiated/ionized raw materials and that this product was not irradiated/ionized.

D-ALPHA TOCOPHERYL ACETATE (700 UI-G) (VIT E NAT)

ANTI-DOPING

As per the list of 2016 of the World Anti-Doping Agency, the ingredient is not a doping substance or a combination of doping substances. The ingredient does not contain any doping substance. The ingredient does not result from a doping substance.

NANOMATERIALS

According to the definition of 'Nanomaterials' of the UE Regulation No 1169/2011 of the European Parliament and of the Council of 25th of October 2011 (amending Commission Regulation (EC) 1924/2006 and Commission Regulation (EC) 1925/2006) on the provision of food information to consumers, we hereby attest that no nanomaterials are used in the manufacture, in the formulation nor in the packaging materials of this product.

D-ALPHA TOCOPHERYL ACETATE (700 UI-G) (VIT E NAT)

NUTRITIONAL INFORMATION

Approximative Nutritional Information of this product is as follows:

TYPICAL NUTRITIONAL INFORMATION

(for 100g of product)

Calories from fat (Kcal) 85

Calories from saturated fat (Kcal) 13.8

Total fat (%,g) 9.6

Saturated fatty acids* (%,g) 1.5

Monounsaturated fatty acids* (%,g) 2.5

Polyunsaturated fatty acids* (%,g) 5.4

Trans fat (%,g) < 1

Total Carbohydrates (%,g) 0

Dietary fiber (%,g) 0

Sugar (%,g) 0

Protein (%,g) 0

Moisture (%,g) <5

Cholesterol by GC (mg/100g) <1

Sodium (mg/100g) < 0.3

Potassium (mg/100g) < 0.1

Total Vitamin E (%) (as d-alpha tocopheryl acetate) 51.4

*calculated as triglyceride

D-ALPHA TOCOPHERYL ACETATE (700 UI-G) (VIT E NAT)

RESIDUE LIMITS (GELATIN, COLLAGEN)

not applicable

HYGIENE OF FOODSTUFFS

We herewith certify that this product complies with Commission Regulation (EC) No. 852/2004 of the European Parliament on the hygiene of foodstuffs and all its amendments.

FULL TRACEABILITY

This product complies with the full traceability criteria as mentioned in Commission Regulation (EC) No. 178/2002 of the European Parliament and of the Council of 28th of January 2002, laying down the general principles and requirements of food law, establishing the European food safety authority and laying down procedures in matters of food safety.

D-ALPHA TOCOPHERYL ACETATE (700 UI-G) (VIT E NAT)

FACTORY ACCREDITATIONS/CERTIFICATION	ONS
ISO 9001	Yes
ISO 22000	No
HACCP	Yes
GMP	Yes
FAMI QS	No
Other accreditations/certifications	FSSC:22000, IP, Halal, Kosher

D-ALPHA TOCOPHERYL ACETATE (700 UI-G) (VIT E NAT)

NON ANIMAL TESTING

We certify that the product has not been tested on animals.



VITAMIN B3, NICOTINAMIDE

Product information:

Product name Nicotinamide Chemical formula $C_6H_6N_2O$ Cas number 98-92-0 EP

Test parameter: Specification:

Appearance A white or almost white, crystalline powder or colourless

crystals, freely soluble in water and in anhydrous ethanol,

slightly soluble in methylene chloride

Appearance of solution Solution should be clear and not more intensely coloured

than reference solution BY7

Odour test by NaOH Passes test
Colour test Passes test

Identification (A) Melting point: 128 – 131°C

(B) IR: complies

(C) Thin-Layer chromatography: The principal spot in the chromatogram obtained with the test solution is similar in

position and size to the spot in the chromatogram

obtained with the reference solution

Assay 99 – 101%

Loss on drying Max. 0,5%

Sulfated ash Max. 0,10%

pH 6 – 7,5

Related substances

Unspecified impurity Max. 0,1%
Total impurity Max. 0,2%
Heavy metals (as Pb) Max. 20 ppm

Additional information:

Shelf life 5 years

Store in its original, unopened packaging in a cool,

dry and well-ventilated place

Packaging 25kg bags
Application Nutritional

Vitamine A Acetate 500 000 UI/g Vitamin A acetate 500 000 IU/g Code: 2203032

SPECIFICATION DATA SHEET / FICHE TECHNIQUE

Manufacturing	
Product name	Vitamin A Acetate
Molecular formula	C22H32O2
Composition	Vitamin A Acetate- Starch sodium octenyl succinate, Arabic gum, DL-α-Tocopherol, Silicon dioxide, Maltodextrin, Vitamin C
Compliance	USP/ EP
Origin	Synthetic
Properties	Specifications
Organoleptic	
Appearance	Pale yellow powder
Physical and chemical	,
Identification	The retention time of the principal peak conforms to which in the reference solution
Loss on drying	≤ 5%
Particle size	≥ 90% pass through #80 mesh
Assay	
Vitamin A Acetate content	≥ 500 000 IU/g HPLC
Microbiological*	
Total plate count	≤ 1 000 CFU/g
Yeast and moild	≤ 100 CFU/g
Enterobacteriae	≤ 10 CFU/g
E.coli	Absent (10g)
Salmonella	Absent (25g)
Staphylococcus aureus	Absent (10g)
Contaminants*	
Lead	≤ 2 ppm
Arsenic	≤ 2 ppm
Mercury	≤ 0.1 ppm
Cadmium	≤ 1 ppm
Shelf Life	2 years
Allergens	Absence

Vitamine A Acetate 500 000 UI/g Vitamin A acetate 500 000 IU/g Code: 2203032

ATTESTATION ALLERGENES / ALLERGENS STATEMENT

Au regard du règlement 1169/2011 et de ses modifiants, veuillez trouver ci-dessous la liste des allergènes dans le produit ci-dessus :

Absence d'allergènes	 Présence d'allergènes (préciser ci-dessous)
----------------------	---

SOURCE D'ALLERGENES	Présence dan	s la préparation
Céréales contenant du gluten et produits à base de ces céréales	□OUI	⊠ NON
Crustacés et produits à base de crustacés	□OUI	⊠ NON
Œufs et produits à base d'œufs	□OUI	⊠ NON
Poissons et produits à base de poissons	□OUI	⊠ NON
Arachides et produits à base d'arachides	□OUI	⊠ NON
Soja et produits à base de soja	□OUI	⊠ NON
Lait et produits à base de lait (y compris le lactose)	□OUI	⊠ NON
Fruits à coque et produits à base de ces fruits	□OUI	⊠ NON
Moutarde et produits à base de moutarde	□OUI	⊠ NON
Céleri et produits à base de céleri	□OUI	⊠ NON
Graines de sésame et produits à base de graines de sésame	□OUI	⊠ NON
Lupin et produits à base de lupin	□OUI	⊠ NON
Mollusques et produits à base de mollusques	□OUI	⊠ NON
Anhydride sulfureux et sulfites en concentration de plus de 10 mg / litre exprimées en SO ₂	□OUI	⊠ NON

In accordance with Regulation (EU) 1169/2011 and its modifications, please find below the list of allergens in the product named above:

X	Without allergens	Alleraens	(specifi	helow
	without allerdens	Alleluella	CODECITO	DEIOW

ALLERGEN SOURCE	Presence in the prepara	tion
Cereals containing gluten or derived products	□YES ⊠ NO	
Crustaceans and derived products	□YES ⊠ NO	
Eggs and derived products	□YES ⊠ NO	
Fishes and derived products	□YES ⊠ NO	
Peanuts and derived products	□YES ⊠ NO	
Soybeans and derived products	□YES ⊠ NO	
Milk and derived products (including lactose)	□YES ⊠ NO	
Tree nuts and derived products	□YES ⊠ NO	
Mustard and derived products	□YES ⊠ NO	
Celery and derived products	□YES ⊠ NO	
Sesame seeds and derived products	□YES ⊠ NO	
Lupins and products with lupin	□YES ⊠ NO	
Mollusks and products with mollusks	□YES ⊠ NO	
The Sulphur Dioxide and Sulphites (E220 to E228) are lower than 10 mg/kg	□YES ⊠ NO	

Vitamine A Acetate 500 000 UI/g Vitamin A acetate 500 000 IU/g Code: 2203032

ATTESTATION DE NON IONISATION / NON IONIZATION STATEMENT

Le produit cité ci-dessus n'a pas été soumis à des rayonnements ionisants conformément aux Directives 1999/2/CE et 1999/3/CE.

According to Directives 1999/2/EC and 1999/3/EC, we confirm that the product named above has not been subjected to ionizing radiation.

ATTESTATION OGM / GMO STATEMENT

Conformément aux Règlements 2001/18/CE, 1829/2003/CE et 1830/2003/CE, nous déclarons que le produit ci-dessus n'est pas génétiquement modifié puisqu'il :

- Ne provient pas de substances génétiquement modifiées
- Ne contient pas de supports ou d'additifs génétiquement modifiés.

According to Regulations 2001/18/EC, 1829/2003/EC and 1830/2003/EC and its modifications, we declare that the product above is not genetically modified as it:

- Does not come from genetically modified raw material
- Does not contain any carrier or additive coming from genetically modified organisms.

ATTESTATION NANOMATERIAUX / NANOMATERIALS STATEMENT

Le produit cité ci-dessus est conforme au Règlement n°1169/2011 du Parlement européen et du conseil du 25 octobre 2011 sur la fourniture d'information sur les aliments pour les consommateurs.

Le produit contient-il des nanotechnologies telles que définies dans le Règlement n°1169/2011? Oui □ Non ⊠

The product named above is in compliance with Regulation n°1169/2011 of the European parliament and the council of October the 25th 2011, on the provision of food information to consumers.

Does the product contain nanotechnology as defined by Regulation RCE n°1169/2011 Yes □ No ⋈

ATTESTATION EST-ESB / TSE-BSE STATEMENT

Le produit cité ci-dessus est conforme à la réglementation européenne relative au risque d'Encéphalopathie Spongiforme Transmissible (EST).

The product named above is in compliance with the European legislation relative to the risk of Transmissible Spongiform Encephalopathy (TSE).

STATUT VEGAN / VEGAN STATUS

Le produit cité ci-dessus est vegan et ne contient aucun produit animal (incluant produits laitiers, œufs, miel...). The product named above is vegan. It doesn't contain any animal products (including dairy products, eggs, honey...).

Scientific name	Biotin
CAS N°	58-85-5
Molecular formula	$C_{10}H_{16}N_2O_3S$
Continent of origin	Europe
Origin	Synthetic
Natural from	/
Melting point:	229-232°C
Ar	NALYTICAL DATA
Appearance	White to off-white crystalline powder
Identification	Match with reference IR spectrum
Assay	97.5-101%
Individual impurity	NMT 1.0%
Total impurities	NMT 2.0%
Loss on drying	NMT 0.1%
Sulphated ash	NMT 0.1%
Specific rotation	+89 - +93°
	NLT 50% <10μm
Particle size	NMT 90% <20μm
	NLT 98% <30µm
STORAGE	CONDITION – SHELF LIFE
Type of packaging	Not Specified
Packaging	Suitable for food industry
Storage conditions	Store at temperature below 25°C, in a well closed bag away from moisture and direct sun light
Shelf life	3 years if stored in accordance with recommendations
Batch size	Batch size depends of our suppliers and of customer order (kg to tons)
Stability data	Not Specified

Vitamin B8 – Biotin – VITB8001

PRODUCT IMPURITIES*

Conta	minants and residues	Ana	alysis	Not tested	Content- limit	Reference
Conta	minants and residues	Each batch	Control plan	tes N	Content- mint	and/or method
	Lead (Pb)	\boxtimes			<3ppm	USP/EP
Heavy metals	Cadmium (Cd)	\boxtimes			<1ppm	USP/EP
He	Mercury (Hg)	\boxtimes			<0.1ppm	USP/EP
	Arsenic (As)	\boxtimes			<2ppm	USP/EP
0	Total plate count	\boxtimes			<20 000 cfu/g	
ntr	Yeast & moulds	\boxtimes			<200 cfu/g	
Microbiological control	Bile tolerant gram – bacteria (enterobacteria)	\boxtimes			<100 cfu/g	
log	E. coli	\boxtimes			Negative/g	
bio	Salmonella	\boxtimes			Negative/25g	
cro	Staphylococcus aureus				Negative/g	
Σ	Others				7/2	
	Residual solvent			\boxtimes	-	
	Pesticides			\boxtimes	<mrl< td=""><td></td></mrl<>	
	Aflatoxin B1				:=	
	Sum of B1, B2, G1, G2				1.5	
	Ochratoxin A				*=	
	Dioxins and PCBs		\boxtimes		NMT 0.75 ng/kg	Dir. 2002/70/EC
2	Benz(o)apyrene				<10ppb	
minants control	Sum of PAH (benzo(a)pyrene, benz(a)anthracene, benzo(b)fluoranthene and chrysene)				<50ppb	
Contam	Melamine and its structural analogues				-	-
W al a	lodine (algae)			\boxtimes	-	
	Microcystin (algae)			\boxtimes	ie.	-
	Other contaminants :					
	Pyrrolizidine alkaloids			\boxtimes	On customer's request	=
	3-MCPD (3-monochloro- propanol-1,2-diol)			\boxtimes	.=	

^{*}Information given by our supplier on the date of updating of data sheet

Vitamin B8 – Biotin – VITB8001

ATTESTATIONS - STATEMENTS

GMOs FREE S	STATEMENT	
According to the manufacturer's declaration and to a Regulation (EC) No. 1829/2003 of the European Pa on genetically modified food and feed - Regulation (EC) No. 1830/2003 of the European Pa concerning tractability and labelling on GMOs and tracked GMOs and amending Directive 2001/18/EC	arliament and of the Coun arliament and of the Coun	cil of 22 September 2003 cil of 22 September 2003
GMOs FREE	⊠ Yes	□ No
If no, name and quantity (%):		
NON-IRRADIATI	ON STATEMEN	Т
According to the manufacturer's declaration and to t the European Parliament and of the Council:	he European Directive 199	99/2/EC and 1999/3/EC of
NON-IRRADIATION	⊠ Yes	□ No
BSE/TSE FREE	STATEMENT	
According to the manufacturer's declaration:		
BSE/TSE FREE	⊠ Yes	□ No
NANOMATERIALS	FREE STATEMEI	NT
According to the manufacturer's declaration and to Parliament and of the Council of 25 October 2011:	the EU Regulation No 11	69/2011 of the European
NANOMATERIALS FREE	⊠ Yes	□ No

If no, name and quantity (%):

Vitamin B8 - Biotin - VITB8001

ALLERGENS AND CROSS CONTAMINATION STATEMENT

According to the manufacturer's declaration and to the EU Regulation No 1169/2011 of the European Parliament and of the Council of 25 October 2011, this product is:

ALLERGENS FREE		⊠ Yes □	□ No	(if no sec	e below)
	Present in the product (Yes/No)	Quantity (ppm) In case of presence	prod	uction sit tion line (allergen on the e or on the same On Natural site and lier site)
Cereals containing gluten, namely: wheat (such as spelt and khorasan wheat), rye, barley, oats or their hybridised strains, and products thereof (1)	(***)	an.	⊠ Yes	□ No	□ Not applicable
Crustaceans and products thereof	(414)	***	⊠ Yes	□ No	\square Not applicable
Eggs and products thereof	***		imes Yes	\square No	\square Not applicable
Fish and products thereof (2)	***	***		□ No	\square Not applicable
Peanuts and products thereof	***	3444	\square Yes	⊠ No	\square Not applicable
Soybeans and products thereof (3)	(***)	1111		□ No	\square Not applicable
Milk and products thereof (including lactose) (4)	***	***	imes Yes	\square No	\square Not applicable
Nuts, namely: almonds (Amygdalus communis L.), hazelnuts (Corylus avellana), walnuts (Juglans regia), cashews (Anacardium occidentale), pecan nuts (Carya illinoinensis (Wangenh.) K. Koch), Brazil nuts (Bertholletia excelsa), pistachio nuts (Pistacia vera), macadamia or Queensland nuts (Macadamia ternifolia), and products thereof (5)	2777.6	-177	⊠ Yes	□No	□ Not applicable
Celery and products thereof	***	***	⊠ Yes	□ No	\square Not applicable
Mustard and products thereof	***	***		□ No	\square Not applicable
Sesame seeds and products thereof	***	***		□ №	\square Not applicable
Sulphur dioxide and sulphites at concentrations of more than 10 mg/kg or 10 mg/litre in terms of the total SO2 (6)	Sees	See	⊠ Yes	□ No	□ Not applicable
Lupin and products thereof	2444	33.0	\square Yes	\boxtimes No	\square Not applicable
Molluscs and products thereof	***	***		□ No	\square Not applicable

The risk of cross contamination on Natural site and supplier site is avoided by cleaning procedures.

¹⁾ Exceptions: Wheat based glucose syrups including dextrose / Wheat based maltodextrins / Glucose syrups based on barley /Cereals used for making alcoholic distillates including ethyl alcohol of agricultural origin

⁽²⁾ Exceptions: fish gelatin used as carrier for vitamin or carotenoid preparation / fish gelatin or Isinglass used as fining agent in beer and wine

⁽³⁾ Exceptions: Fully refined soybean oil and fat / natural mixed tocopherols (E306), natural D-alpha tocopherol acetate, and natural D-alpha tocopherol succinate from soybean sources / vegetable oils derived phytosterols and phytosterol esters from soybean sources / plant stanol ester produced from vegetable oil sterols from soybean sources

⁽⁴⁾ Exceptions: whey used for making alcoholic distillates including ethyl alcohol of agricultural origin / lactitol

⁽⁵⁾ Exceptions: nuts used for making alcoholic distillates including ethyl alcohol of agricultural origin

⁽⁶⁾ In terms of the total SO2 which are to be calculated for products as pronosed ready for consumption or as reconstituted according to the instructions of the manufacturers

Vitamin B8 - Biotin - VITB8001

EUROPEAN FOOD REGULATIONS

EUROPEAN REGULATIONS		COMP	LIANT
ontaminants (Regulation 1881/2006/EC)	⊠ Yes	□ No	□ Not applicable
eavy metals (Regulation 629/2008/EC)	⊠ Yes	□ No	□ Not applicable
esticides (Regulation 396/2005/EC)	□ Yes	□ No	⊠ Not applicable
xtraction solvents (Directive 2009/32/EC)	□ Yes	□ No	⊠ Not applicabl
AH and benzoapyrene (Regulation 2015/1933/EC)	☐ Yes	□ No	⊠ Not applicable
FOOD GRA cording to the manufacturer's declaration, this product	is:	□ No	⊠ Not applicable
FOOD GRA cording to the manufacturer's declaration, this product	DE is:		
cording to the manufacturer's declaration, this product	IDE	□ No	
FOOD GRA coording to the manufacturer's declaration, this product	DE is:		Not applicable□ Not specify⋈ Not analyzed
FOOD GRA coording to the manufacturer's declaration, this product STATEMENTS Food grade	is:	□ No	□ Not specify
FOOD GRA coording to the manufacturer's declaration, this product STATEMENT: Food grade Gluten free (<20mg/kg - Regulation 828/2014)	ADE is: S	□ No	□ Not specify ☑ Not analyze
FOOD GRA coording to the manufacturer's declaration, this product STATEMENTS Food grade Gluten free (<20mg/kg - Regulation 828/2014) Suitable for vegetarians	is: ✓ Yes ✓ Yes ✓ Yes	□ No □ No	□ Not specify ☑ Not analyze □ Not specify

According to the manufacturer's declaration and to the European regulations Directive 2003/15/CE and 2003/80/CE:

NON-ANIMAL TESTING	☐ Yes	□ No	☑ Not specify

Manganese gluconate

rodu				

Product Name	Manganese gluconate	
INCI Name	bis(D-gluconato-O1,O2)manganese	
EINECS Number	Number 229-350-4	
CAS Code	6485-39-8	

Product:	Manganese Gluconate Dihydrate USP28 Powder
Appearance	Light pink coloured powder
Identification	
A	To pass test
В	To pass test
Assay (on dry basis)	98.0-102.0%
Water	6-9%
Chloride	0.05% max.
Sulfate	0.2% max.
Arsenic	2mg/kg max.
Lead	3mg/kg max.
Mercury	0.1mg/kg max.
Cadmium	1mg/kg max.
Heavy metals	20ppm max.
Reducing substances	1.0% max.
Organic volatile impurities	Meets requirements
Manganese content(as dihydrate)	11.19%-11.64%
Particle size Thru 70 mesh	
Microbial Limits	
Total aerobic count	10000cfu/g max.
Total molds-yeasts	100cfu/g max.
Salmonella	Absent/10gram
Enterobacteriaceaes	Absent/1gram
E.Coli	Absent/1gram

Manganese gluconate

GENERAL PRODUCT INFORMATION

70
0.81 g/ml
0.97 g/ml
80g/L at DEG C
Synthetic
Mineral
Yes
e)? Yes
Yes

Pharmacopoeias that product meets	USP/FCC
Drug indication	N/A
Melting Point	204 Deg C

Manganese gluconate

BSE/TSE

The product does not contain and is not derived from Specified Risk Material as defined in Commission Directive European and it is conform to the EU legislation relating to the risk of Transmission of Spongiform Encephalopathy (TSE).

GMC

Following EU regulations have been published in the Official Journal of the European Union:

- Regulation (EC) No.1829/2003 of the European Parliament and the Council of 22nd September 2003 on genetically modified food and feed
- Regulation (EC) No.1830/2003 of the European Parliament and the Council of 22nd September 2003 concerning tractability and labeling on genetically modified organisms and tractability of food and feed products produced from genetically modified organisms and amending Directive 2001/18/EC.

This product has not been genetically modified so there is no obligation of GMO labeling as defined by the above mentioned regulations.

NON IRRADIATION/NON IONIZED

According to directives 1992/2/EC and 1999/3/EC, We confirm is not made from irradiated/ionized raw materials or was irradiated/ionized the product.

Manganese gluconate

PESTICIDES

To our actual knowledge of production process, raw materials and equipment used potential pesticide residues in the above mentioned product comply with the European legislation on pesticide residues, esp. Regulation (EC)None. 396/2005. However, we do no test for pesticide residues on a routine basis.

RESIDUAL SOLVENTS

The product meets with:

- DIRECTIVE 2009/32/EC OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 23 April 2009 on the approximation of the laws of the Member States on extraction solvents used in the production of foodstuffs and food ingredients
- DIRECTIVE 2010/59/EU of 26 August 2010 amending Directive 2009/32/EC of the European Parliament and of the Council on the approximation of the laws of the Member States on extraction solvents used in the production of foodstuffs and food ingredients.

CONTAMINANTS

The product is in accordance and meets with the commission regulation (EC) No.629/2008 of 2 July 2008 amending regulation (EC) No.1881/2006 setting maximum levels for certain contaminants in foodstuffs.

PCBS and MELAMINE

The product is in accordance and meets with the commission regulation (EU) No.594/2012 of 5 July 2012 amending regulation (EC) No.1881/2006 as regards the maximum levels of the contaminants Ochratoxin a, non dioxin-like PCBS and melamine in foodstuffs.

ANTI-DOPING

As per the list of 2012 of the world anti-doping agency the ingredient is not a doping substance and not a combination of doping substances. The ingredient does not contain any doping substance. The ingredient does not result from a doping substance.

As per the list of 2016 of the world anti-doping agency the ingredient is not a doping substance and not a combination of doping substances. The ingredient does not contain any doping substance. The ingredient does not result from a doping substance.

NANOMATERIALS

According to the the definition of 'nanomaterials' of the UE REGULATION (EU) No 1169/2011 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 25 October 2011 on the provision of food information to consumers, we hereby attest that no nanomaterials are used in the formulation nor in the packaging materials of our product.

Manganese gluconate

ALLERGENS

In accordance with the Directive 2003/89/EC and Council amending Directive 2000/13/EC and taking Directive 2005/26/EC, Directive 2005/63/EC and Directive 2006/142/EC of 22.12.2006 into consideration. We confirm the following information:

Note: Does the raw material CONTAIN or is it DERIVED from or could it be CROSS-CONTAMINATED by any of the following.

Cereals containing gluten (i.e. wheat, rye, barley, oats, spelt, kamut or their hybridised strains) and products thereof	No
Crustaceans and products thereof	No
Eggs and products thereof	No
Fish and products thereof	No
Peanuts and products thereof	No
Soybeans and products thereof	No
Milk and products thereof (including lactose)	No
Nuts (i.e. Almond, Hazelnut, Walnut, Cashew, Pecan nut, Brazil nut, Pistachio nut, Macadamia nut and Queensland nut and products thereof)	No
Celery and products thereof	No
Mustard and products thereof	No
Sesame seeds and products thereof	No
Sulphur dioxide and sulphite at concentrations of more than 10 mg/kg or 10 mg/l expressed as SO2	No
Lupine and products thereof	No
Molluscs and products thereof	No

Produit: POTASSIUM IODURE

01 - IDENTIFICATION DE LA SUBSTANCE / PREPARATION ET DE LA SOCIETE

DENOMINATION POTASSIUM IODURE

CODE ARTICLE 5 010 043

NOM DU FOURNISSEUR COOPERATION PHARMACEUTIQUE FRANCAISE

Place Lucien Auvert 77020 MELUN Cedex

Nom INCI Potassium iodide

 Numéro CAS
 7681-11-0

 Numéro EINECS
 231-659-4

Formule brute KI

Masse molaire 166,01 g/mol

02 - SPECIFICATIONS

REFERENTIEL DE CONTROLE Pharmacopée Européenne en vigueur

Caractères Poudre blanche ou sensiblement blanche ou cristaux incolores

Solubilité dans l'eau Très soluble

Solubilité dans la glycérine Facilement soluble

Solubilité dans l'alcool Soluble

Identification par contenant Positive

Iodures Positive

Potassium Positive

Aspect de la solution Limpide et incolore

Alcalinité Maximum 0,5 ml HCl 0,01 M

Iodates Conforme

Sulfates Maximum 150 ppm

Thiosulfates Conforme

Fer Maximum 20 ppm

Métaux lourds Maximum 10 ppm

Perte à la dessiccation Maximum 1,0 %

Teneur en iodure de potassium sur produit sec 99,0 à 100,5 %

03 - CONDITIONNEMENTS

250 g Code article: 1 603 610

1 kg Code article: 1 603 800

25 kg Code article: 1 603 820

04 - AUTRES INFORMATIONS

LISTE DES MODIFICATIONS SUCCESSIVES Version 1 du 21/09/07 : création.

Version 2 du 10/01/08 : modification § 2.

AUTRES INFORMATIONS Les renseignements contenus dans cette fiche sont basés sur l'état de

nos connaissances relatives au produit concerné à la date de mise à

jour.

Les informations relatives au stockage et à la sécurité sont

mentionnées dans la fiche de données de sécurité.

L'attention des utilisateurs est attirée sur les risques éventuellement encourus lorsqu'un produit est utilisé à d'autres usages que ceux pour

lesquels il est destiné.

Fin du document

Chromium picolinate

	SPECIFICATION D	DATA		0
Nom chimique / Chemical Name	m chimique / Chemical Name Chromium picolinate			
Numéro CAS / CAS number	14639-25-9			N N N
Numéro EINECS / EINECS number) cr
Masse Molaire / Molecular mass	418.33			0,000
Formule brute / Molecular Formula	C ₁₈ H ₁₂ N ₃ O ₆ Cr			EN TOO
Conformité / Compliance	USP	727		
Utilisation possible / Use	Nutraceutique		Nutraceuti	cal
Apparence / Appearance	Poudre		Powder	
Odeur et couleur / Odor and Color	Rouge à rose		Red to pink	(
Granulométrie / Particle size	NLT 90% # 60 mesh	·	· · · · · · · · · · · · · · · · · · ·	
Solubilité / Solubility	Insoluble dans l'eau et	l'alcool	Insoluble in	n water/alcohol
Densité / Bulk density	NLT 0.45g/mL	- 10		
Excipient / Carrier	Aucun		None	
	Méthode d'analyses	Fréquence d		Spécifications
	Analysis Method *	Frequency o	f analysis	Specifications
Teneur/ Assay	USP	Chaque lot / E	ach batch	98-102% on dried basis
Perte à la dessication/ Loss on drying	USP	Chaque lot / E	ach batch	≤ 4.0%
Cendres sulfuriques /Sulphated ashes				< 0.2%
рН				58 (0.5% solution)
Pouvoir rotatoire / Specific rotation				process constant who can proceed to restrict the Manager who we are
Point de fusion / Melting point				253-245°C
Identification / Identification	A. IR spectrum	Chaque lot / E	ach hatch	Comply with standard
identification/ identification	B. Chemical test	Chaque lot / E		AS per USP
Autres / Other	D. Chemical test	Situação ioti, s		As per osi
Chloride	USP	Chaque lot / E	ach hatch	≤ 0.06%
Sulphate	USP		ot / Each batch ≤ 0.20%	
Chromium	A	Chaque lot / Each batch		12.18-12.68%
	WI .			22.20 22.00//
Métaux lourds totaux / Total heavy	1			
metals				
Arsenic / Arsenic	In house	Chaque lot / E	ach batch	≤1 ppm
Plomb / Lead	In house	Chaque lot / E	ach batch	≤3 ppm
Cadmium / Cadmium	In house	Chaque lot / E	ach batch	≤1 ppm
Mercure / Mercury	In house	Chaque lot / E		≤ 0.1 ppm
	- I was a second and a second a			
Flore totale / Total Plate count (CFU/g)	In house	Chaque lot / E	ach batch	≤1000
Levures et moisissures /Yeast & Mold (CFU/g)	In house	Chaque lot / E	ach batch	≤ 100
E. Coli	In house	Chaque lot / E	ach batch	Absent/g
Salmonella	In house	Chaque lot / E	U 45 H 47 45	Absent/g
	1 10 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2		1 7 7 7 6

	Chi	romium picolinate			
storage	conditions (T°)	room temperature			
Conditio	nnement / Packaging	25 kg hmhdpe DRUMS			
Durée de conservation / Shelf life 5 ans / 5 years					
	GENETICALLY	MODIFIED ORGANISMS – GM	0		
		egulations EC/1829/2003 & EC/1830/200			
	g to the EU regulations concerning the traceab products produced from GMO (Regulation N				
1.	> Contains ingredient(s) from Maize and,	or Soya origin	Yes 🗌	No 🖾	
		onsists of or is produced from authorized . This presence is adventitious or technically	Yes 🖂	No 🗆	
	> is GMO or contains or consists of GMO	S	Yes 🗌	No 🖾	
	> is produced from GMOs or contains ing	gredients produced from GMOs	Yes 🗌	No 🖾	
2.	 Contains ingredient(s) from other pote cotton) 	ntial GMO-risk raw material(s) (rice, beet,	Yes 🗌	No 🖾	
	If Yes, please specify :				
	o ingredient :				
	o vegetable origin :				
	o geographic origin :				
		onsists of or is produced from evaluated . This presence is adventitious or technically	Yes 🛚	No 🗆	
3.	For information about processing aids,				
	is produced or contains ingredient(s) pro	oduced	\	N- M	
	with GMOs or Genetically Modified Microorganisms processing aids (bacteria, yeast) or derived from GMOs or GMMs (including enzymes)				
	If yes, please specify :				
25 S					
	irm that the ingredient supplied is compliant 003/EC and N°1830/2003/EC where applicable	in all respects with the Regulations	Yes 🛛	No 🗌	

Chromium picolinate

INGREDIENTS

An ingredient is defined in Directive 2000/13/EC on labelling: "ingredient means any substance including additives, <u>used in the production</u> or preparation of a foodstuff and still present in the finished product although possibly in a modified form" i.e. all raw materials to produce the goods.

Please list all ingredients:

%	Ingredients/material composition
.00	Chromium picolinate
	emonium pieemate

SUBSTANCES OR PRODUCTS CAUSING ALLERGIES OR INTOLERANCES

According to Regulation (EU) N°1169/2011 as regards of ingredients present in foodstuffs

The modification of the 2000/13 European directive about the labelling aims at the obligation of labelling clearly the allergen below used in food products production and still present in the product, even in a modified form, including additives and processing aids.

	Direct incorporation as a base raw material or in a derivative form (don't forget the carriers, the additives and the processing aids)		CROSS CONTAMINATION					
			Presence on the production line		Presence on the production workshop		Presence on the production factory	
	Yes (precise nature)	No	Yes (precise the nature)	No	Yes (precise the nature)	No	Yes (precise the nature)	No
Cereals containing gluten(1) and products thereof	13706	x		x		×		x
Crustaceans and products thereof		х		х		х		х
Eggs and products thereof		×		х		×		×
Fish and products thereof (2)		х		x		х		х
Peanuts and products thereof		x		х		х		х
Soybeans and products thereof (3)		×		x		×		×
Milk and products thereof (including lactose)(4)		х		х		×	(2)	x
Nuts (5) or products thereof		×		×		×		×
Celery and products thereof		х		х		x		х
Mustard and products thereof		×		×		×		×
Sesame seeds and product thereof		х		х		×		х
Sulphur dioxide and sulphites at concentrations of more than 10mg/kg or 10 mg/l expressed as SO2		×		x		×		×
Lupines and products thereof		×		x		×		х
Mollusc and product thereof		х		х		х		Х
Latex		×		×		x		×

⁽¹⁾ Cereals which contain gluten (i.e. wheat, rye, barley, oats, spelt, kamut or their hybridised strains) except: wheat-based glucose syrups including dextrose, wheat-based maltodextrins, glucose syrups based on barley, cereals used for making distillates or ethyl alcohol of agricultural origin for spirit drinks and other alcoholic beverages:

⁽²⁾ Except: fish gelatine used as carrier for vitamin or carotenoid preparations, fish gelatine or Isinglass used as fining agent in beer and wine;

⁽³⁾ Except fully refined soybean oil and fat, natural mixed tocopheols (E306), natural D-alpha tocopherol, natural D-alpha tocopherol succinate from soybean sources; vegetable oils derived phytosterols and phytosterol esters from soybean sources; plant stanol ester producted from vegetable oil sterols from soybean sources;

⁽⁴⁾ Except when used for making distillates or ethyl alcohol of agricultural origin for spirit drinks and other alcoholic beverages, lactitol;

Chromium picolinate

(5) almond (Amydalus communis L.) hazelnuts (Corylus avellana), walnut (Juglans regia), cashew (Anacardium occidentale), pecan nuts (Carya illinaiesis), brazil nut (Bertholletia excelsa), pistachio nut (Pistacia vera), macadamia nut and queensland nut (Macadamia terniflora) and products thereof, except nuts used for making distillates or ethyl alcohol of agricultural origin for spirit drinks and other alcoholic bereverages;

ALLERGENS & FOOD INTOLERANCE

	Direct incorporation as a base raw material or in a derivative form (don't forget the carriers, the additives and the processing aids)			CROSS CONTAMINATION				
			Presence on the production line		Presence on the production workshop		Presence on the production factory	
	Yes* (precise nature)	No	Yes* (precise the nature)	No	Yes* (precise the nature)	No	Yes* (precise the nature)	No
Gluten	P 200	×	98	х	200	×		х
Yeasts and hydrolysates, cardamom, vanilla, cinnamon, cocoa, leguminosaes; benzoic acid, benzoate, parabens, phenylalanine, glutamate, coriander and product thereof.		×		x		x		x
Bovine, porcine, ovine, fish and by- products		×		х		x		×
Exotics fruits (kiwi, banana, mango,) and products thereof		×	2	х		x		×
Strawberry and others berries		х	43	х		×		х
Azo-colorants	6	x		x		x		х
Latex		х		х		х		х
Others:								

*	ıf	Yes	
	ш	162	

 could you please send us your risk assessment and/or your Allergens p

Does this raw material have been made with nanotechnology?

Does the product contain nanotechnology as defined by the French decree n°2012-232?

- could you please sella as your risk assessment and/c	your Allergens policy:				
- Procedure in place to prevent or control cross-conta	cts: specify the procedures:				
☐ Validated cleaning procedure (s) ☐ Raw material segregation / specific label					
☐ Specific production scheduling ☐ Dedicated equipment / production areas					
Other: please specify					
Attach documents describing these procedures.					
- Please specify bellow the name of the source of alle	ergen and its quantity:				
NOI	N IRRADIATION				
According to	UE Directive N°1999/2/CE				
Has this raw material or one of its ingredients been irr	adiated or ionised?	YES	NO⊠		
NA	NOMATERIAL				
According to F	Regulation (EU) N°1169/2011				
Does this raw material contain or can be considered a	s engineered nanomaterial*2	VES	NOM		

'engineered nanomaterial' means any intentionally produced material that has one or more dimensions of the order of 100 nm or less or that is composed of discrete functional parts, either internally or at the surface, many of which have one or more dimensions of the order of 100 nm or less, including structures, agglomerates or aggregates, which may have a size above the order of 100 nm but retain properties that are characteristic of the nanoscale.

YES

Copper gluconate

Product SPECIFICATION

Product Name	Copper gluconate
INCI Name	copper di-D-gluconate
EINECS Number	208-408-2
CAS Code	527-09-3

The product meets OR exceeds: USP30	
Product:	Copper Gluconate Anhydrous USP30 and EU Regulation Powder
Identification	
A	To pass test
В	To pass test
Assay	98.0 - 102.0%
Sulfate	0.05% max.
Chloride	0.07% max.
Arsenic (As)	1 mg/kg max.
Lead (Pb)	3 mg/kg max.
Mercury (Hg)	0.1 mg/kg max.
Cadmium (Cd)	1 mg/kg max.
Reducing substance	1.0% max.
Copper content	13.72 ~ 14.28%
Particle size thru a 70 mesh	
Microbial Limits	
Total aerobic count	10000 cfu/gram max.
Total molds-yeast	100 cfu/gram max.
Salmonella	Absent/25 gram
Enterobacteriaceaes	Absent/1 gram
Staphylococcus aureus	Absent/1 gram
Listeria monocytogenes	100cfu/gram

Copper gluconate

GENERAL PRODUCT INFORMATION	
Mesh of the product	70
Taped denisty	0.47g/ml
Bulk density	0.73g/ml
Water solubility	45.3-49g/100mL at 25 DEG C
Origin (Synthetis, Fermentation)	Synthetic
Origin (Animal, Vegetal, etc.)	Mineral
Is the product and their ingredients FREE of GLUTEN?	Yes
Product is suitable for Vegans	Yes
Product is suitable for Vegatarians	Yes
Product is free of fructose	Yes
Is this product suitable for food?	Yes

Yes

Pharmacopoeias that product meets	USP FCC	
Drug indication	N/A	
Melting Point	167 Deg C	

Has been sold this product in the EU before May 1997?

Can be the product used in baby food (According to Directive 2006/141/CE)? Yes

Copper gluconate

TECHNICAL PRODUCT INFORMATION	
Name of the product	Copper Gluconate
IUPAC name	Copper Gluconate
ATC Code	N/A
Molecular Formula	C12H22CuO14
Stereochemistry	N/A
Molecular mass	453.84 g/mol
Raw materials used and their function	Gluconic Acid and Cu Carbonate belong to food additive
Frequency of analysis of each contaminant?	Each Batch
Frequency of analysis of microbiology?	Each Batch, Each Batch
Frequency of analysis of each heavy metals?	Each Batch, Each Batch

Copper gluconate

BSE/TSI

The product does not contain and is not derived from Specified Risk Material as defined in Commission Directive European and it is conform to the EU legislation relating to the risk of Transmission of Spongiform Encephalopathy (TSE).

GMO

Following EU regulations have been published in the Official Journal of the European Union:

- Regulation (EC) No.1829/2003 of the European Parliament and the Council of 22nd September 2003 on genetically modified food and feed
- Regulation (EC) No.1830/2003 of the European Parliament and the Council of 22nd September 2003 concerning tractability and labeling on genetically modified organisms and tractability of food and feed products produced from genetically modified organisms and amending Directive 2001/18/EC.

This product has not been genetically modified so there is no obligation of GMO labeling as defined by the above mentioned regulations.

NON IRRADIATION/NON IONIZED

According to directives 1992/2/EC and 1999/3/EC, We confirm is not made from irradiated/ionized raw materials or was irradiated/ionized the product.

Copper gluconate

PESTICIDES

To our actual knowledge of production process, raw materials and equipment used potential pesticide residues in the above mentioned product comply with the European legislation on pesticide residues, esp. Regulation (EC)None. 396/2005. However, we do no test for pesticide residues on a routine basis.

RESIDUAL SOLVENTS

The product meets with:

- DIRECTIVE 2009/32/EC OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 23 April 2009 on the approximation of the laws of the Member States on extraction solvents used in the production of foodstuffs and food ingredients
- DIRECTIVE 2010/59/EU of 26 August 2010 amending Directive 2009/32/EC of the European Parliament and of the Council on the approximation of the laws of the Member States on extraction solvents used in the production of foodstuffs and food ingredients.

CONTAMINANTS

The product is in accordance and meets with the commission regulation (EC) No.629/2008 of 2 July 2008 amending regulation (EC) No.1881/2006 setting maximum levels for certain contaminants in foodstuffs.

PCBS and MELAMINE

The product is in accordance and meets with the commission regulation (EU) No.594/2012 of 5 July 2012 amending regulation (EC) No.1881/2006 as regards the maximum levels of the contaminants Ochratoxin a, non dioxin-like PCBS and melamine in foodstuffs.

ANTI-DOPING

As per the list of 2012 of the world anti-doping agency the ingredient is not a doping substance and not a combination of doping substances. The ingredient does not contain any doping substance. The ingredient does not result from a doping substance.

As per the list of 2016 of the world anti-doping agency the ingredient is not a doping substance and not a combination of doping substances. The ingredient does not contain any doping substance. The ingredient does not result from a doping substance.

NANOMATERIALS

According to the the definition of 'nanomaterials' of the UE REGULATION (EU) No 1169/2011 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 25 October 2011 on the provision of food information to consumers, we hereby attest that no nanomaterials are used in the formulation nor in the packaging materials of our product.

Copper gluconate

ALLERGENS

In accordance with the Directive 2003/89/EC and Council amending Directive 2000/13/EC and taking Directive 2005/26/EC, Directive 2005/63/EC and Directive 2006/142/EC of 22.12.2006 into consideration. We confirm the following information:

Note: Does the raw material CONTAIN or is it DERIVED from or could it be CROSS-CONTAMINATED by any of the following.

Cereals containing gluten (i.e. wheat, rye, barley, oats, spelt, kamut or their hybridised strains) and products thereof	No
Crustaceans and products thereof	No
Eggs and products thereof	No
Fish and products thereof	No
Peanuts and products thereof	No
Soybeans and products thereof	No
Milk and products thereof (including lactose)	No
Nuts (i.e. Almond, Hazelnut, Walnut, Cashew, Pecan nut, Brazil nut, Pistachio nut, Macadamia nut and Queensland nut and products thereof)	No
Celery and products thereof	No
Mustard and products thereof	No
Sesame seeds and products thereof	No
Sulphur dioxide and sulphite at concentrations of more than 10 mg/kg or 10 mg/l expressed as SO2	No
Lupine and products thereof	No
Molluscs and products thereof	No

Sélénomethionine L Poudre Se 0.5% - Origine synthétique Selenomethionine L Powder Se 0.5% - Synthetic origin Code: 1602163

SPECIFICATION DATA SHEET - FICHE TECHNIQUE

Manufacturing	
Manufacturing Product name	2-amino-4-methylselanyl-butanoic acid
Molecular formula	C5H11NO2Se
	196.106 g/mol
Molecular weight CAS N°	3211-76-5
Origin	Synthetic Solicium contrarate > 80%
Carrier	Calcium carbonate > 90%
Properties	Specifications
Organoleptic	
Appearance	White crystalline powder to off white powder
Chemical	
Se content	≥ 0.5% (5000ppm)
Physical	
Loss on drying	≤ 10.0%
Particle size	≥ 90% pass through #40 mesh
Microbiological	
Total plate count	≤ 3000 CFU/g
Yeast & Mold	≤ 300 CFU/g
E.coli	Absence
Salmonella	Absence
Staphylococcus aureus	Absence
Heavy metals	
Lead	≤ 3ppm
Arsenic	≤ 1ppm
Cadmium	≤ 1ppm
Mercury	≤ 0.1ppm
Shelf life	2 years

^{*}According to a control plan

Remarks: To be stored in original tightly closed package away from moisture and sunlight

Abbreviations: ND: not determined / NA: not applicable

Sélénomethionine L Poudre Se 0.5% - Origine synthétique Selenomethionine L Powder Se 0.5% - Synthetic origin Code: 1602163

ATTESTATION ALLERGENES / ALLERGENS STATEMENT

Au regard du Règlement (UE)	1169/2011	ainsi que ses modifications,	certifie que:
-----------------------------	-----------	------------------------------	---------------

SOURCE D'ALLERGENES	Présence dan	Présence dans la préparation		
Céréales contenant du gluten et produits à base de ces céréales	□OUI	⊠ NON		
Crustacés et produits à base de crustacés	□oui	⊠ NON		
Œufs et produits à base d'œufs	□oui	⊠ NON		
Poissons et produits à base de poissons	□oui	⊠ NON		
Arachides et produits à base d'arachides	□OUI	⊠ NON		
Soja et produits à base de soja	□oui	⊠ NON		
Lait et produits à base de lait (y compris le lactose)	□oui	⊠ NON		
Fruits à coque et produits à base de ces fruits	□OUI	⊠ NON		
Moutarde et produits à base de moutarde	□OUI	⊠ NON		
Céleri et produits à base de céleri	□OUI	⊠ NON		
Graines de sésame et produits à base de graines de sésame	□OUI	⊠ NON		
Lupin et produits à base de lupin	□oui	⊠ NON		
Mollusques et produits à base de mollusques	□OUI	⊠ NON		
Anhydride sulfureux et sulfites en concentration de plus de 10 mg / litre exprimées en SO ₂	□oui	⊠ NON		

In accordance with Regulation (EU) 1169/2011 and its modifications, certifies that:

Without allergens ■ Without allergens	☐ Allergens (specify below

ALLERGEN SOURCE	Presence in th	Presence in the preparation		
Cereals containing gluten or derived products	□YES	⊠ NO		
Crustaceans and derived products	□YES	⊠ NO		
Eggs and derived products	□YES	⊠ NO		
Fishes and derived products	□YES	⊠ NO		
Peanuts and derived products	□YES	⊠ NO		
Soybeans and derived products	□YES	⊠ NO		
Milk and derived products (including lactose)	□YES	⊠ NO		
Tree nuts and derived products	□YES	⊠ NO		
Mustard and derived products	□YES	⊠ NO		
Celery and derived products	□YES	⊠ NO		
Sesame seeds and derived products	□YES	⊠ NO		
Lupins and products with lupin	□YES	⊠ NO		
Mollusks and products with mollusks	□YES	⊠ NO		
The Sulphur Dioxide and Sulphites (E220 to E228) are lower than 10 mg/kg	□YES	⊠ NO		

Sélénomethionine L Poudre Se 0.5% - Origine synthétique Selenomethionine L Powder Se 0.5% - Synthetic origin Code: 1602163

ATTESTATION DE NON IRRADIATION / NON IONIZED STATEMENT

Le produit et les matières qui le composent n'ont pas été soumis à des rayonnements ionisants conformément aux Directives 1999/2/CE et 1999/3/CE.

According to Directives 1999/2/EC and 1999/3/EC, we confirm that the product and the ingredients used in it have not been subjected to ionizing radiation.

ATTESTATION OGM / GMO STATEMENT

Conformément aux Règlements 2001/18/CE, 1829/2003/CE et 1830/2003/CE, nous déclarons que le produit ci-dessus n'est pas génétiquement modifié puisqu'il :

- Ne provient pas de substances génétiquement modifiées
- Ne contient pas de supports ou d'additifs génétiquement modifiés.

According to Regulations 2001/18/EC, 1829/2003/EC and 1830/2003/EC and its modifications, we declare that the product above is not genetically modified as it:

- Does not come from genetically modified raw material
- Does not contain any carrier or additive coming from genetically modified organisms.

ATTESTATION NANOMATERIAUX / NANOMATERIALS STATEMENT

Nous certifions que ce produit est conforme au Règlement n°1169/2011 du Parlement européen et du conseil du 25 octobre 2011 sur la fourniture d'information sur les aliments pour les consommateurs.

We certify that this product is in compliance with Regulation n°1169/2011 of the European parliament and the council of October the 25th 2011, on the provision of food information to consumers.

ATTESTATION EST-ESB / TSE-BSE STATEMENT

Nous certifions que notre produit est conforme à la réglementation européenne relative au risque d'Encéphalopathie Spongiforme Transmissible (EST).

We certify that this product is conform to the European legislation relative to the risk of Transmissible Spongiform Encephalopathy (TSE).

ATTESTATION CONFORMITE EMBALLAGES / PACKAGING STATEMENT

Nous certifions par la présente, que le conditionnement de ce produit satisfait aux législations suivantes :

- Règlement n°10/2011/CE de la Commission du 14 janvier 2011 concernant les matériaux et objets en matière plastique destinés à entrer en contact avec les denrées alimentaires
- Règlement n° 1935/2004/CE du Parlement européen et du conseil du 27 octobre 2004 concernant les matériaux et objets destinés à entrer en contact avec des denrées alimentaires et abrogeant les Directives 80/590/CEE et 89/109/CEE

Hereby, we certify that the packaging used complies with:

 Commission Regulation n°10/2011/EC dated January the 14th 2011, on plastic materials and articles intended to come into contact with food and its modifications

TECHNICAL DATA SHEET

	ILUII	ITTICAL	. DAIA SI			
		PRODU	UCT NAME			
	VITAMINE K2 1% MENAQUINONE-7 MK-7					
		REF	ERENCE			
**************************************		VIT	TK2.1.01			
		SUBSTANCE	E INFORMATION			
Grade	Food Grade		CAS N°	2124-57-4		
			Carrier (%)	None		
Shelf life	2 years					
ACTIVI	INGREDIENTS /	SUBSTANCE	S TO CONTROL			
Assay		NLT 1% Men	aquinone-7 MK-7	HPLC		
	ANALY1	TICAL DATA				
Appearance			Yellow powde	er		
Melting Point			53-55°C			
	CONTA	AMINANTS				
Heavy metals			NMT 20ppm			
Lead (Pb)			NMT 2ppm			
Arsenic (As)			NMT 2ppm			
Cadmium(Cd)			NMT 0,3ppm	ri		
Mercury (Hg)		NMT 0,3ppm				
	MICRO	OBIOLOGY				
Total aerobic microbia	l		NMT 20 000cfu	/g		
Tot. yeast and mould			NMT 200cfu/g	g		
Salmonella			Negative/25g	3		
E.Coli			Negative/g			
	CERT	TIFICATE				
Allergens			Allergen free	8		
GMO		Non OGM				
Irradiation			Non Irradiatio	n		
	PACKAGING	AND STORA	AGE			
Packing		М	lust be suitable for foc	od industry		
Storage		In a cool a	and dry place, away fro	om direct sun ligh		

FICHE TECHNIQUE

AMIDON DE RIZ BIO origine EUROPE -SAC 20KG

Description of product

"Organic Rice starch" is a native starch extracted from "ORGANIC" rice of 100% EU origin.

Extraction from selected organic rice is carried out by mechanical action and using pure water; drying does not change product properties which is marketed in the form of white light powder, insoluble in fresh water, with a neutral taste and flavour. The Organic certification is issued by Italian Authority recognized with Ministerial Ordinance.

The product is available

- . in bulk;
- . in 20 Kg multiple paper bags; the pallets 100×120 cm. or EPAL 80×120 cm (wrapped with polyethylene film);
- . Big Bags in polypropylene of different sizes (90x90x160 90x90x180) available on pallets 100x120.

Tariff Heading

11081910

Producer Code

8282B: Organic Rice Starch EU

8282B20: Organic Rice Starch EU in 20kg bags 8282BBB: Organic Rice Starch EU in bigbags Distributeur certifié par Ecocert France SAS (FR-BIO-01)

Storage

keep product in a dry and cool place, at room temperature, in its sealed packaging.

Transport

- . bulk product: by clean and dry trucks, with the necessary health permissions and HACCP certification if product is destined for human consumption;
- . product in bags: by dry, clean and odourless trucks with a waterproof cover.

Intended use

- . food (cakes, breakfast cereals, baby food);
- . pharmaceutical
- . cosmetic

Shelf life

3 (three) years from the date of production, under the conditions described in "storage".

Product Specification

*	Value guaranteed	Method	CoA	
Chemical-Physical Parameter				
Moisture	14,0 % max	IR	X	
рН	6,0 - 7,5	20% sol.	X	
Ashes	0,60 % d.s. max	IR	X	
Proteins (N*5,7)	1,00 % d.s. max	IR	X	
Sieving Test				
>50µm	20,0 % max	Sieving		
>500µm	0,1 % max	Sieving		
Viscosity Brabender (Method: 40 g d.s. in water 500 ml /d.h.: 350 cmg)	70.000 + 5	Brabender		
Starting Gel Point	70,0 °C ± 5	380100000000000000000000000000000000000		
Brabender end viscosity	300 BU	Brabender		
Microbiological Parameter				
Bacterial Total Aerobic Count (CFU/g.)	10.000 max	3M Petrifilm	X	
Enterobacteria (CFU/g.)	10 max	3M Petrifilm	X	
Moulds (CFU/g.)	500 max	3M Petrifilm	X	
Yeasts (CFU/q.)	100 max	3M Petrifilm	X	

Nous ne nous engageons pas à fournir un produit conforme aux applications baby food et cosmétiques Sauf indication contraire dans l'intitulé , nos produits sont de grade ALIMENTAIRE/FOOD.

AMIDON DE RIZ BIO origine EUROPE -SAC 20KG

HACCP Declaration

This product is manufactured in conformity to the EU Regulation 852/2004 on the food hygiene.

Nutritional Information

Parameter	Medium value on 100 g
Moisture	12,0
Energy (kj)	1467
Energy (kcal)	351
Proteins	0,50
Carbohydrates	86,1
Of which sugars	<0,1
Fat	0,50
Of which satured	0,15
Fiber	0,50
Salt	0,10
Ashes	0,40

Allergens declaration

3									
Allergens	Present in the product production line			Present on site		Cross contamination risk in the product			
	YES	8	YES	NO	YES	No.	YES	8	
Cereals containing gluten (*) and products thereof		х		х	х			х	
Crustaceans and crustaceans based products		X		x		x		х	
Eggs and egg-based		X		X		X		X	
Fish and fish products		X		X		X		X	
Peanuts and peanut products		X		X		X		X	
Soy and soy products		X		X		X		X	
Milk and milk-based products (including lactose)		×		x		x		х	
Nuts (**) and products thereof		X		X		X		X	
Celery and celery-based products		X		X		X		X	
Mustard and mustard-based products		x		х		×		х	
Sesame seeds and products made from sesame seeds		х		x		×		х	
Sulphur dioxide and sulphites at concentrations above 10 mg / kg or 10 mg / litre expressed as SO2		x		х	X			х	
Lupine and lupine-based products		X		X		X		X	
Molluscs and molluscs based products		X		х		х		×	

Nous ne nous engageons pas à fournir un produit conforme aux applications baby food et cosmétiques Sauf indication contraire dans l'intitulé, nos produits sont de grade ALIMENTAIRE/FOOD.

^{(*):} Wheat, rye, barley, oats, spelled, kamut or their hybridised strains
(**): Almond (Amygdalus communis L.), hazelnuts (Corylus avellana), walnuts (Junglans director), cashew (Anacardium occidentale), pecan
[Carya illinoiesis (Wangehn) K.Koch], brazil nuts (Brazil nut), pistachio (Pistacia vera), Queensland nuts (macadamia ternifolia)

AMIDON DE RIZ BIO origine EUROPE -SAC 20KG

GMO Declaration

- it originates from the workmanship of rice or broken rice;
- it doesn't contain GMO;
- ✓ it is gotten by raw materials NON GMO;
- during the production any GMO ingredient and/or additive has not been employed.

Contaminants declaration

We guarantee that every batch is conforming to the EU Reg. 1881/06 for the presence of heavy metals residual (Pb, Cd) and micotoxins.

Regarding the contaminants above mentioned we perform in external laboratory at least 1 analysis on 1 batches of ORGANIC RICE STARCH EU

We declare that our products are not treated with radiation.

Certification

The product is certified Kosher and Halal

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