



RAW MATERIAL DATA SHEET

PrincipHYAL Line

PRODUCT IDENTIFICATION					
(I) Supplier name	ROELMI HPC SRL Legal office: Via Celeste Milani, 24/26 – 21040 Origgio (VA) - Italy T. +39 02 3351 0150 F. +39 02 3354 9210				
	This RMDS refers to all the PrincipHYAL products, except to the "full spectrum" line (Cube ³ , Difference, Aurora, Signal 10)				
(II) EU/US INCI name					
	EU INCI name	US INCI nam	e		
	Sodium	Sodium			
	Hyaluronate	Hyaluronate	2		
(III) Regulatory information	Empirical Formula: (C14H20NO11Na)n Chemical name: Sodium Hyaluronate				
	Common n	name	CAS n.	EC n.	Composition
	Sodium Hyalı	uronate	9067-32-7	(polymer)	100%
	US and European Pharmacopoeia: Not applicable Classification & labelling according to EU Chemical regulation: this product is not subjected to labelling as it is not a dangerous substance (regulation 1272/2008). Transport: not regulated				
	Tariff code: 39139000	0			
(IV) Functions	Humectant				
. ,	Skin conditioning				
(V) Geographical origin	PrincipHYAL Line is produced in our production plant located in China				
REGULATORY STATUS					

ROELMI HPC srl

REA: VA-361664

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Capitale sociale Euro 120.000 i.v.











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(VI) Chemical Inventory	Component(s) listed in EU Inventory: yes (Sodium Hyaluronate)				
position	Component(s) listed in IECSC (China): yes (Sodium Hyaluronate)				
	Component(s) listed in TSCA (USA): no				
	Component(s) listed in NZIoC (New Zeland): yes (Sodium Hyaluronate)				
	Component(s) listed in TCSI (Taiwan): no				
	Component(s) listed in ECNS (Japan): no				
	Component(s) listed in KECI (Korea): no				
	Component(s) listed in PICCS (Philippines): no				
	Component(s) listed in AICS (Australia): yes (Sodium Hyaluronate)				
	Component(s) listed in DSL (Canada): yes (Sodium Hyaluronate)				
	Preposition 65 (California): not listed				
(VII) Cosmetic Regulatory	No restrictions in Europe.				
Status	Other countries: PrincipHYAL Line is accepted in most other legislative areas.				
	Please refer to your local legislation of interest.				
	Restriction or limitation for personal care applications: no				
	Canadian cosmetic ingredient hotlist: no				
	Component listed in Cosmetic China IECIC: yes				
(VIII) Cosmetic Regulatory	Regulatory Conformity: PrincipHYAL Line is in line with Cosmetics Regulation				
Conformity					
	substances listed in Annex II and Annex III of cosmetics regulation.				
	CMR: PrincipHYAL Line does not contain any substances intentionally added and				
	classified as Carcinogenic, Mutagenic or Reprotoxic substances, based on the				
	classification criteria of the Regulation (EC) 1272/2008 (CLP).				
	Allergens: We certify that PrincipHYAL Line complies with Regulation (EC) 1223/2009				
	regarding the presence of the 26 substances identified as allergenic in cosmetics. Due to				
	the nature of the materials used for our productions and to bibliographic data, it can be				
	stated that these allergens are absent.				
	TSE/BSE: We confirm that PrincipHYAL Line does not contain any animal raw material,				
	therefore it is free from BSE/TSE transmitting agents				
	Animal testing: We declare that no animal testing was made by Roelmi HPC & C. srl for				
	cosmetic use accordingly to Regulation (EC) 1223/2009.				
	<u>lonization</u> : the product is not subjected to ionization process				
	REACH: According to Article 2(9), polymers are exempted from the provisions on				
	regulation of Title II of REACH.				
	PrincipHYAL Line meets the criteria set for a polymer defined in Article 3(5).				
	Its monomers can be considered as non-isolated intermediates, for which REACH				
	regulation is not applicable according to Article 2(1).				
	Based on the available data, the criteria for exemption from REACH registration are				
	fulfilled for Sodium Hyaluronate (CAS: 9067-32-7).				

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(IX) Toxicology Toxicological information

Ecotoxicity Biodegradability information

Tests for cosmetic safety evaluation (sponsored by the company)

Skin irritation test: NOT IRRITANT (method: MTT/ %Tested: 0.5% & 0.1% / Date: April 2009)

Ocular irritation test: NOT IRRITANT (method: Red Blood Cell Test/ %Tested: 1% & 10% /

Date: April 2009)

Skin sensitization test: NON SENSITIZING (method: Repeated Patch test / %Tested: Pure

100% / Date: April 2009)

Mutagenicity test: Not available

Photo toxicity test/ Photo allergencity: test Not available

Bibliographic data*

Eye irritation (Draize test): non irritant

Skin irritation: non irritant (Japanese rabbits and Hartley guinea pig)

Skin sensitization): non sensitizing (Guinea Pigs - Comparison between Sensitization of

Non-crosslinked (SupArtz tes) and Crosslinked (Synvisc test) Hyaluronic Acid)

Reproductive and Developmental Toxicity:

NOAEL (Rats): 60 mg/Kg (Source: Rooster combs/ Treatment: subcutaneous injection

from day 17 of pregnancy to day 20 after parturition)

NOAEL (Female rats): 64 mg/kg (Source: Bacteria/Treatment: subcutaneous injection from day 7 today 17 of pregnancy)

*CIR- Final Report of the Safety Assessment of Hyaluronic Acid, Potassium Hyaluronate, and Sodium Hyaluronate. (July/August 2009)

Following our bibliographic research on our raw material, to date no scientific data and/or reference has been found which could be interpreted as a sign of toxicity. This lead us to assume it is not toxic when used at normal dose.

Ecological & Biodegradability information

Biodegradability: expected to be biodegradable

and possible traces of Heavy metals: see SDS contaminants

(X) Unavoidable impurities | SVHC (as per REACH regulation): absence

Pesticides: absence

GMO: absence

Food allergens (as per Regulation (EC) no. 1169/2011) and gluten: absence (no starting

materials containing food allergens or gluten are used in the product).

Residual solvents: yes (maximum content < 0,5% Ethanol)

Polycyclic Aromatic Hydrocarbons: absence

Free amines: absence

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Phthalates: absence

Glycol ethers: absence Formaldehyde: absence Ethylene oxide: absence

Nanomaterials: absence Monomers: absence Proteins: see SDS

VOC (volatile organic components as per European and Swiss laws): maximum content

< 0.5% Ethanol

Other impurities or residues: absence

Absence: based upon data from our starting material suppliers and knowledge of the manufacturing process, we have no reason to believe that these substances are

present.

(XI) Additives | Neutralisers: no.

Preservatives (following annex V of Regulation (EC) NO. 1223/2009): no.

Antioxidants: no. Stabilisers: no. Catalysts: no.

Bleaching agents: no. Chelating agents: no.

Ethanol: yes (maximum content < 0,5% Ethanol*)

*Vegetal origin, main sources: corn, potatoes.

(XII) Ingredients origin | Synthetic: No

Animal: No

Animal Protection (Cites): NA (not applicable)

<u>Vegetable</u>: No

Plant name: NA (not applicable)

Plant Protection (Cites): NA (not applicable)

Palm oil and derivates: No.

GMO: No

Mineral: No

Polymer: yes (Sodium Hyaluronate)

Biotechnological processing: Yes

Component name: Sodium Hyaluronate

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	Strain used for biotechno Components of the ferme	_	n: Streptococcus equi subsp. Zooepidemicus
	Substrate/Organism	Origin	
	Peptones	Wheat	
	Glucose	Corn starch	
	Yeast extract	Yeast	
			erials and processing aids used in ntain or consist of GMO's and they are not
(XIII) Halal/Kosher	We confirm that the prod these substances are use Roelmi HPC will communi	d during manufactu	01
(XIV) Vegan/vegetarian	PrincipHYAL Line does no	t contain any anima	l origin ingredients or animal derivatives.
PRODUCT INFORMATION			
(XV) Properties	Please refer to marketing		
	MICROBIOLOGICAL STABI		urfacturing process and to the low
	Due to the renning steps	included in the Mar	ufacturing process and to the low

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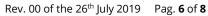
availability of water in the finished product we can guarantee the microbiological

specifications of the SDS for any batch (absence of common pathogens).







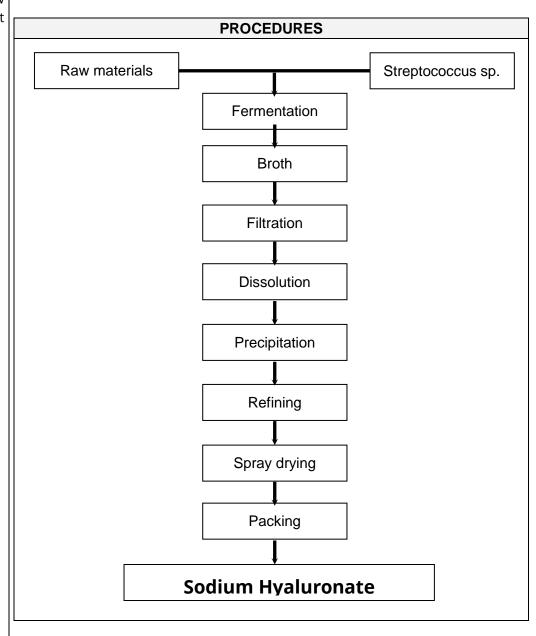




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(XVI) Manufacturing flow chart



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(XVII) Specification	SDS available	
(XVIII) Material safety data	MSDS or equivalent statement available	
sheet		
(XIX) Personal Care	Particular instructions for the use of PrincipHYAL Line in cosmetic applications:	
Applications and	Solubility:	
formulation	Slowly but fully dissolves in water to give viscous, clear to slightly opalescent and colourless solutions.	
	Temperature:	
	The ingredient is not sensitive to usual temperature range. It can be heated up to 80° C for 1 to few hours before starting/accelerating degradation process.	
	Incorporation:	
	Prepare a pre-solution of the powder (e.g.: at 1%) in water, at ambient temperature. Mix under mechanical agitation for approximately 40 minutes, until totally dispersed. Include then the solution:	
	- either in aqueous phase before mixing	
	- either in formed emulsion during cooling phase	
	Compatibility:	
	Sodium Hyaluronate is a large molecule polyanion polysaccharide. During dissolution, avoid cation emulsifiers or cation preservatives to keep a stable and transparent formula.	
	pH range: The ingredient is stable at cosmetic pH: its rheological attitude is perfectly maintained in a pH range of 2.8 – 12.0.	
	For further information, let's note that around pH 2.5, a thermo-reversible gel-like behaviour can be registered, attributed to an interaction mechanism. At pH lower < 2.5, the polymer is re-solubilized again, and the solution-gel transition is pH-reversible. At higher pH (> 12.5), a transition to a random coil configuration can occur. This phenomenon is due to the –OH groups dissociation in alkaline conditions, and a consequent reduction of –OH bonds that control the physiological stiffness of the hyaluronan molecule.	
	Other: This ingredient is very hygroscopic, and should be maintained in closed packaging away	
	from humidity. It is preferable to avoid turbo mixer.	
	Formulation information:	
	The typical dosage is: 0.1 – 2.0 %	
	Its use is recommendable for the following cosmetic products:	
	- Emulsion, cream, gel, lotion, serum, mask	
	- Skin care, body-care, make-up	
	- Face, eye-contour, baby % specific	

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	Usual finished product applications are diverse: - Anti-aging formulas - Lifting & boosting solutions
	- Multi-function formulas - Short-term hydration
	- Sagging or inelastic skins - Mature & young skins
(XX) Efficacy test	No test for efficacy
(XXI) Storage and handling	Keep the product in the original container well closed in fresh and dry place, repaired from humidity, light or heat sources.
(XXII) Shelf Life	36 months from the production date** **24 months from production date if the molecular weight is greater than 1800kDa.

THE INFORMATION CONTAINED IN THIS DOCUMENT ARE BASED ON OUR CURRENT SCIENTIFIC KNOWLEDGE AND DOES NOT DISCHARGE THE USER FROM THE CONTROL OF THE PRODUCT BEFORE THE USE. THE INFORMATION ARE CAREFUL AND RELIABLE BUT IT SHOULD NOT BE TAKEN AS EXPRESSING OR IMPLYING ANY WARRANTY CONCERNING THE PRODUCT CHARACTERISTICS. THE USER, UNDER HIS OWN RESPONSIBILITY, MUST RESPECT THE RULES AND LAWS IN FORCE. ROELMI HPC SRL DOES NOT TAKE LIABILITY FOR ANY DAMAGE TO PEOPLE OR PROPERTY FOR IMPROPER USE AND DECLINES ANY RESPONSABILITY IN CASE THE USE OF THE PRODUCT INFORMATION MIGHT TAKE TO PATENT INFRINGEMENT OF THIRD PARTIES.

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