ADDITIONAL TECHNICAL DOCUMENTATION

COLLAGENE IDROLIZZATO PE

002838 | Rev. 1 | 14/01/2026 | EN

English rendering of the supplied raw-material documentation (kollagen.pdf). Source information and selections are retained; this is not a newly issued manufacturer certificate.

INCI	Aqua, Hydrolyzed Collagen
INCI USA	Water, Hydrolyzed Collagen
CAS	7732-18-5, 92113-31-0/9015-54-7
EINECS / ELINCS	231-791-2, 295-635-5
Composition	Aqua: > 50,0 % Hydrolyzed Collagen: 25,0–50,0 %

* The stated ranges refer to the quantities of raw materials used during production. The quantity of plant used (expressed as fresh or rehydrated plant) is conventionally indicated as '…extract'.

Product origin
[X] Animal: Hydrolyzed Collagen. Animal-synthetic (from bovine skin).
[ ] Vegetable
[ ] Mineral
[X] Synthetic: Phenoxyethanol, Ethylhexylglycerin
[ ] From technological processes (fermentation, hydrolysis …)
[X] Other: Tetrasodium glutamate diacetate: natural-synthetic.

Restrictions listed in CITES Annexes: [ ] Yes [X] No
If yes, specify: _____

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Information on the manufacturing process — additives / impurities

Substance group	NO*	YES / INCI	CAS	%
Amines / nitrosamines	X			
Antioxidants	X			
Preservatives		Phenoxyethanol	122-99-6	0,90
Additives	X	Ethylhexylglycerin Tetrasodium glutamate diacetate	70445-33-9 51981-21-6	0,10 0,094
1,4-dioxane	X			
Formaldehyde and formaldehyde releasers	X			
Phthalates	X			
Gluten**	X			
Lactose**	X			
Latex	X			
Palm oil and derivatives	X			
Ethylene oxide	X			
Residual solvents	X			
Other (specify)	X			

* For impurities listed as absent: To the best of our current knowledge of the raw materials used and the manufacturing process, these substances are not expected to be present, except through accidental cross-contamination. No analytical testing has been conducted in support of this statement.

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** Allergenic substances: The extract concerned does not contain intentionally added allergenic ingredients listed in Annex II to Regulation (EU) 1169/2011. It is stored in a facility where allergenic ingredients under that regulation are present. Procedures have been implemented to avoid cross-contamination. However, botanical products cannot be guaranteed entirely free of potentially allergenic substances because environmental and/or accidental contamination may occur in the field.

To the best of our current knowledge and according to raw-material suppliers' statements, the following limits are expected to be respected:

HEAVY METALS: Compliant with the requirements of Regulation (EC) No 853/2004 and subsequent amendments.

PESTICIDES: Compliant with Regulation (EC) No 396/2005 and subsequent amendments.

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Additional regulatory information

ANIMAL TESTING — Article 18 of Regulation (EC) No 1223/2009 of the European Parliament and of the Council of 30 November 2009 on cosmetic products.

Compliance (not tested on animals for cosmetic purposes after 11 March 2009): [X] Yes [ ] No

CMR***: If the product contains substances classified as CMR categories 1A, 1B or 2 under Part 3 of Annex VI to Regulation (EC) No 1272/2008 (CLP), at concentrations equal to or greater than the substance’s specific or generic concentration limit, these are listed in the safety data sheet in accordance with Article 11 of Regulation (EC) No 1272/2008.

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NANOMATERIALS — Article 2.1.k, Chapter I, of Regulation (EC) No 1223/2009 of the European Parliament and of the Council of 30 November 2009 on cosmetic products.

Nanomaterial: An insoluble or biopersistent and intentionally manufactured material with one or more external dimensions, or an internal structure, on the scale of 1–100 nm.

Is the product classified as a nanomaterial under this definition? [ ] Yes [X] No

COSMETIC ALLERGENS — Commission Regulation (EU) 2023/1545 of 26 July 2023 amending Regulation (EC) No 1223/2009 of the European Parliament and of the Council as regards labelling of fragrance allergens in cosmetic products.

The product:
[ ] Contains
[X] Is not expected to contain***

*** No analytical testing has been conducted.

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BSE / TSE — Regulation (EC) No 220/2009 of the European Parliament and of the Council.

The product:
[ ] Is of non-bovine origin and therefore exempt from BSE/TSE risk.
[X] Is of bovine origin.

Only in the latter case, specify:
Animal species: Bovine
Organs used: Skin
Country of origin: _____

The manufacturer declares that the starting collagen used to produce the solution is free from TSE (Transmissible Spongiform Encephalopathy) and BSE (Bovine Spongiform Encephalopathy) risks.

GMO — Regulations (EC) No 1829/2003 and No 1830/2003 and European Directive 2001/18/EC.

The product:
[ ] Contains / comes from
[X] Does not contain / does not come from

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Treatment with ionising radiation

The product:
[ ] Has been treated
[X] Has not been treated

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