

Technical Data Sheet

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Version 2 from 13/04/2011

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1.	INFORMATION ON INGREDIENT	
1.1	Trade Name	dermofeel® PA-12
1.2	Manufacturer	
1.3	Supplier	Dr. Straetmans GmbH Merkurring 60 -62 D-22143 Hamburg, Germany Tel.: +49 - (0)40 - 66 93 56 0 Fax: +49 - (0)40 - 66 93 56 10 E-Mail: info@dr-straetmans.de
1.4	Ingredient Category	Chelating agent
1.5	Chemical Description	myo-Inositol hexakis dihydrogenphosphate sodium salt
1.6	INCI-Name	Sodium Phytate
1.7	EINECS / ELINCS	238-242-6
1.8	CAS-no.	14306-25-3
1.9	for colors: C.I. Colipa-no.	Not applicable
1.10	Registration Status	EU: registered US: registered Japan: JCIC 559718

2.	MANUFACTURING PROCESS INFORMATION	
2.1	Origin of starting material	Vegetable
2.2	Description of manufacturing process	Extraktion from corn with diluted mineral acid and subsequent ion exchange
2.3	Residues from Manufacturing	
2.3.1	Catalyst	Not to be expected
2.3.2	Solvent	Up to 4% Water
2.3.3	Others	Not to be expected
2.4.	Storage Time	12 months

3.	SPECIFICATION	
3.1	Assay	Min 96%
3.2	Additives	
3.2.1	Preservation, %	None
3.2.2	Antioxidants, %	None
3.2.3	Solvent, %	None
3.2.4	Other Additives, %	None

3.3	Microbiological Specification	
3.3.1	Total Plate Count (cfu/g) Aerobic Germs (cfu/g)	No contamination expected No contamination expected
3.3.2	Pathogene Organisms	No contamination expected
3.4	Impurities	
3.4.1	1,4-Dioxan	Not to be expected
3.4.2	Ethylenoxide	Not to be expected
3.4.3	Residual Solvent	
3.4.4	Monomers	Not to be expected
3.4.5	Free Amines	Not to be expected
3.4.6	Nitrosamines	Not to be expected
3.4.7	Heavy Metals as Pb (Ni, Pb, As, Cd, Cu, Sb, Cr, Hg)	< 30 ppm
3.4.8	Pesticides	Not known
3.4.9	Polyaromatic Hydrocarbons	Not known
3.4.10	Other Impurities	Not known
3.5	Analysis	
3.5.1	Methods Detections Limits	Specification and methods see attachment
3.5.2	Analysis of Impurities (Methods Detections Limits)	Specification and methods see attachment

4.	TOXICOLOGY	
4.1	Acute Toxicology	LD ₅₀ oral [mg/kg]: Rat ca. 1030 mouse ca. 1030
4.2	Dermal Irritation	Irritating
4.3	Mucous Irritation	Irritating
4.4	Sensitization	Not reported
4.5	Genotoxicity (e.g. Ames-Test)	Not mutagenic
4.6	Percutaneous Permeation	Not known
4.7	Subchronic Toxicity	Not known
4.8	Teratogenicity	None
4.9	Toxicokinetics	Not known
4.10	Additional Toxicitytests	Not known
4.11	Phototoxicity	Not known
4.12	Photosensitization	Not known
4.13	Inhalative Toxicity	Not applicable

5.	HUMAN EXPERIENCE	
5.1	Data on Human Dermal Irritation	At pH-values usually employed in cosmetic products no negative effects of the product has been reported
5.2	Data on Human Sensitization	At pH-values usually employed in cosmetic products no negative effects of the product has been reported

6.	ECOLOGICAL DATA	
6.1	Biological Elimination	Readily biodegradable (OECD 301 A & F)
6.2	Aquatic Toxicity	EC ₅₀ Daphnia Magna (48h): > 100 mg/l (OECD 202)
6.3	Water Endangering Class	1 (self-classification)
6.4	Bioaccumulation Potential	Not known
6.5	Other Information	None

7.	SAFETY DATA SHEET	see attachment
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