

Zschimmer & Schwarz Italiana - 13038 - Tricerro (VC) / ITALY

INFORMAZIONI TOSSICOLOGICHE TOXICOLOGICAL INFORMATION

Revisione n° Revision n° 13

1.	Informazioni generali General information	
1.1	Nome commerciale Trade name	AMPHOTENSID GB 2009 CONC
1.2	Produttore/Fornitore (indirizzo, telefono, fax, contatto) Manufacturer/Supplier (address, phone no., fax no., contact person)	ZSCHIMMER & SCHWARZ ITALIANA Via A. Ariotto 1/C - 13038 Tricerro (VC) Italy Tel: +39 (0)161 808111 Fax: +39 (0)161 801002 e.merlo@zschimmer-schwarz.com
1.3	Categoria della material prima (es. tensioattivo anionico) Raw material category (e.g. anionic surfactant)	Amphoteric surfactant
1.4	Nome chimico Chemical name	Reaction products of 1H-Imidazole-1-ethanol, 4,5-dihydro-,2-(C7-C17 odd-numbered, C17-unsatd. alkyl) derivs. and sodium hydroxide and chloroacetic acid
1.5	Nome INCI (CTFA) Composition INCI (CTFA) name Composition	Disodium Cocoamphodiacetate: average value 39% as active matter, 46% - 53% as dry matter Sodium chloride (facultative in INCI): average value 11% Aqua: to 100%
1.6	N° EC (EINECS-/ELINCS) EC (EINECS/ELINCS) no.	931-291-0 (previous 272-043-5)

1.7	N° CAS CAS no.	68650-39-5
	CAS 110.	
1.8	Registrazioni (es. UE, USA, Giappone) - REACh - Certificazione Registration status (e.g. EU, USA, Japan) - REACh - Certification	TSCA (USA), DSL (Canada), IECSC (China), NZIoC (New Zealand), TCSI (Taiwan), PICCS (Philippines), AICS (Australia) and KECI (Korea). Japanes have recently changed their system, so that publication in the Japanese list of approved ingredients is no longer necessary. Any cosmetic ingredient is now allowed in Japan with no prior approvation. Product is not a biocidal according to Regulation 528/2012. The product is not a phytosanitary according to Regulation 1107/2009. REACh registration n° 01-2119487973-19-XXXX None of substances listed in the "candidate" list of substances of very high concern (SVHC) are contained in the product in a relevant amount. PO 65 (California law): see points 5.6, 5.7 and 5.9. Product is according to the Chapter of Chemical products and substances (Rev. 08, July 2015). ISS code: AUT-10 Toxicological properties are evaluated on the full surfactant (UVCB).

2.	Informazioni sulla produzione Information on production	
2.1	Origine della materia prima (vegetale, animale, sintetica) Origin of starting material (plant, animal, synthetic)	Vegetable (44%), mineral (10%) and synthetic origin (46%). Fatty acids are from vegetable origin. They come mainly from palm kernel oil from Elaeis Guineenis (Malaysia and Indonesia). Aminoethylethanolamine and monochloroacetic acid are synthetic. NaOH is mineral.
2.2	La materia prima deriva da organismi geneticamente modificati (OGM)? Is the starting material derived from	No

	genetically modified organisms (GMO)?	
2.3	Informazioni sul processo di produzione (descrizione generale) Information on production process (general description)	O RCOH + HOCH ₂ CH ₂ -NH-CH ₂ CH ₂ NH ₂ $\xrightarrow{\triangle}$ Fatty acids AEEA ciclyzation NaOH, 2CICH ₂ COOH O CH ₂ COONa RC - NHCH ₂ CH ₂ N - CH ₂ CH ₂ OCH ₂ COONa + NaCI

3.	Additives	
3.1	Conservanti/Biocidi Preservatives/Biocides	Not added and not expected
3.2	Antiossidanti Antioxidants	Not added and not expected
3.3	Solvents Solvents	Water
3.4	Sbiancanti Bleaching agents	Not added and not expected
3.5	Altri Others	Not added and not expected

4.	Specifiche microbiologiche Microbiological specification	
4.1	Conta microbica totale (ufc/g)	less than 10 ufc/g
	Total viable count (colony-forming units/g)	

5. Residui del processo di lavorazione La presenza di tracce delle sostanze elencate in Allegato II del Regolamento No. 1223/2009 (che sostituisce la

Direttiva 76/768/CEE) (incl. CMR cat. 1A, 1B e 2 sostanze contrassegnate con *) deve essere dimostrata come presenza tecnicamente inevitabile lavorando in GMP e deve essere conforme all'Articolo 17 del Regolamento No. 1223/2009.

By-products

The presence of traces of the substances listed in Annex II of Regulation No. 1223/2009 (replaced Directive 76/768/EEC) (incl. cmr cat. 1A, 1B and 2 substances marked with *) shall be allowed provided that such presence is technically unavoidable

5.1	1,4-Diossano * 1,4-Dioxane *	Based on information concerning the raw materials, production process and equipment used it is not likely to be present.
5.2	Ossido di etilene * Ethylene oxide *	Based on information concerning the raw materials, production process and equipment used it is not likely to be present.
5.3	Solventi residui Residual solvents	Based on our actual knowledge of our production process, raw materials and equipment used, no solvent is used in the manufacturing process, only water
5.4	Monomeri residui Residual monomers	Based on information concerning the raw materials, production process and equipment used they are not likely to be present.
5.5	Ammine Amines	Aminoethanolamine: 10 ppm maximum Amidoamine: 0.4% maximum
5.6	Nitrosammine Nitrosamines	Not expected, but because amino groups are present, please avoid to use it with substances able to originate nitrosoamines, that means nitrosating agents (e.g. 2-Bromo-2-Nitropropan-1,3-Diol, NO ₃ , NO ₂) and use proper packaging. In the raw materials (aminoethylethanolamine) and in the product itself, N-Nitrosoamine are below 50 ppb. They are not checked for every batch but in a random way and till now they have always been under detection limits (20 ppb). The product is according to European Cosmetic Legislation. To be preserved in a nitrite-free container.
5.7	Metalli pesanti Heavy metals	Lead (Pb) < 0.25 ppm, Arsenic (As) < 0.25 ppm, Cadmium (Cd) < 0.25 ppm, Chromium (Cr) < 0.50 ppm, Nickel (Ni) < 0.25 ppm, Mercury (Hg) < 0.25 ppm

5.8	Acido monocloroacetico Monochloroacetic acid	600 ppm maximum as sodium monochloroacetate
5.9	Acido dicloroacetico Dichloroacetic acid	30 ppm maximum as sodium dichloroacetate
5.10	Allergens	Based on information concerning the raw materials, production process and equipment used fragrance allergens as of EU Regulation 1223/2009 Annex III, No. 67-92 are not likely to be present. Based on information concerning the raw materials, production process and equipment used food allergens as of EU Directive 2000/13/EC (as amended), Annex IIIa and Regulation (EU) 1169/2001, Annex II are not likely to be present.
5.11	Altri (e.g. CMR) Others (e.g. CMR)	Sodium chloride: 9% - 12% Sodium glycolate: 1.5% maximum Based on information concerning the raw materials, production process and equipment used CMR substances according to Annex VI of the CLP Regulation (EC) 1272/2008 are: aminoethylethanolamine (CMR1B) as impurity technically unavoidable even working in GMP (10 ppm maximum).
6.	Tossicologia Toxicology	In any case toxicological properties are evaluated on the full composition (UVCB substance)
6.1	Informazioni sulla tossicità acuta Information on acute toxicity	LD50 (rats) > 5000 mg/kg (unpublished reports)
6.2	Informazioni sull'irritazione cutanea Information on skin irritation	- From 10% to 12% of active matter on health skin and on skin with lesions = From non irritating to strict irritating (J. of American College of Toxicology, Vol. 9, n° 2, 1990) - 1% active matter, SIDI test = Non irritating (J. of American College of Toxicology, Vol. 9, n° 2, 1990) - 3% in water on health skin and on skin with lesions = Non irritating (Our test n° 1-3-171/3-

75 (1975))

6.3	Informazioni sull'irritazione oculare Information on irritation of the mucous membrane	- Draize test, from 10% to 12% of active matter without rinsing = From light to strict irritant (J. of American College of Toxicology, Vol. 9, n° 2, 1990) - Draize test, from 10% to 12% of active matter with rinsing = From non irritating to slight irritant (J. of American College of Toxicology, Vol. 9, n° 2, 1990)
6.4	Informazioni sulla sensibilizzazione Information on sensitisation potential	- RIPT semiocclusive test, 0.4% in active matter = Not sensitizing (J. of American College of Toxicology, Vol. 9, n° 2, 1990) - RIPT occlusive test, 0.7% in active matter = Not sensitizing (J. of American College of Toxicology, Vol. 9, n° 2, 1990)
6.5	Informazioni sulla genotossicità Information on gene toxicity	Ames test = None mutagenic effects with and without metabolic activation (J. of American College of Toxicology, Vol. 9, n° 2, 1990)
6.6	Informazioni sull'assorbimento percutaneo Information on percutaneous permeation	Not determined
6.7	Altri (e.g. NOAEL) Others (e.g. NOAEL)	General Population Oral DNEL = 0.39 mg/kg bw/day (repeated dose toxicity), AF = 240 (for NOAEL); General Population Inhalation DNEL = 4.06 mg/m³ (repeated dose toxicity), AF = 60 (for NOAEC); General Population Dermal DNEL = 3.85 mg/kg bw/day (repeated dose toxicity), AF = 240 (for NOAEL) Photoxicity: The product doesn't induce photoallergic reactions (J. of American College of Toxicology, Vol. 9, n° 2, 1990)
7.	Ecotossicità	
	Ecology	
7.1	Degradabilità/Eliminazione Degradability/Elimination	Aerobic biodegradability: rapidly biodegradable 84% after 28 d (OECD 301A), 72% after 27 d (OECD 301E), 72% after 28 d (OECD 301F) Anaerobic biodegradability: biodegradable 100% after 28 d (OECD 311)

7.2	Tossicità acquatica acuta Acute aquatic toxicity	- CL50 (fish Oncorhynchus mykiss) = 5.3 mg/l (96h) - CE50 (Daphnia magna) = 8.9 mg/l (48h) - CE50 (Algae Pseudokirchneriella subcapitata) = 16.9 mg/l (72h)
7.3	Altri Others	NOEC: 66 mg/l

8. Informazioni aggiuntive

(Per i dettagli sulle specifiche vedere il bollettino tecnico allegato; per i dettagli sull'etichettatura e la classificazione vedere la scheda di sicurezza allegata.)

Additional information

(For details on specification see enclosed instruction sheet; for details on labelling and classification see enclosed safety data sheet.)

Dichiarazione BSE/TSE

BSE/TSE statement

Dichiarazione test animali

Non-animal testing declaration

Glicol eteri, glicoli, alcoli (non dichiarati nei paragrafi precedenti)

Glycol ethers, glycols, alcohols (not declared in previous paragraphs)

Ftalati, DINP (diisononyl phtalate) Phtalates, DINP (diisononil ftalato)

Parabeni Parabens

Siliconi Silicons The product is not from animal origin. Furthermore it doesn't contain any ingredient of animal origin, it is not produced using ingredients of animal origins and it doesn't come into contact with animal origin ingredients at any stage of its production. It is therefore BSE/TSE free.

ZSCHIMMER & SCHWARZ ITALIANA has never made or commissioned animal tests on this product for cosmetic purpose.

Based on information concerning the raw materials, production process and equipment used they are not likely to be present.

Based on information concerning the raw materials, production process and equipment used phthalates listed in EU Regulation 1223/2009 Annex II are not likely to be present.

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Glutine Gluten

Formaldeide

Formaldehyde (Formol)

VOC

VOC compounds

Fitofarmaci

Plant protection products

APEO, cloroparaffine, composti organici alogenati

APEOs, chloroparaffines, AOX

Mercaptani

Mercaptanes

Melamine Melamine

Lattosio Lactose

Aflatossine/Micotossine Aflatoxines/Mycotoxines

Lattice Latex

Nitrati e Nitriti Nitrates and Nitrites

Amine aromatiche Aromatic amines

Coloranti azoici Azo dyes Based on information concerning the raw materials, production process and equipment used it is not likely to be present.

Based on information concerning the raw materials, production process and equipment used formaldehyde is not likely to be present.

The product doesn't contain any of the substances that are classified as VOC according to "Ordonnance sur taxe d'incitation sur les composes organiques volatils (OCOV) du 12 novembre 1997" or according to Directive 2004/42/EC.

Based on information concerning the raw materials, production process and equipment used plant protection products are not likely to be present.

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The product doesn't contain natural latex and that natural latex is not used/produced in any step of the production process.

Based on information concerning the raw materials, production process and equipment used they are not likely to be present.

Based on information concerning the raw materials, production process and equipment used aromatic amines are not likely to be present.

Based on information concerning the raw materials, production process and equipment used azo dyes are not likely to be present.

Ormoni, antibiotici, steroidi e altri ingredienti (naturali o chimici) dannosi per le funzioni del corpo umano

Hormones, antibiotics, steroids and other ingredients (natural or chemical) dangerous for the body functionality

Pork free Pork free

PBT/vPvB PBT/vPvB

Materiale radioattivo Radioactive material

Nanomateriali **Nanomaterials**

Idrocarburi Policiclici Aromatici Plycyclic Aromatic Hydrocarbons (HAP)

Clorobenzeni/Clorofenoli Chlorobenzenes/Chlorophenols

Ritardanti di fiamma Flame retardants

Grado cosmetico Cosmetic grade

Certificato Kosher Kosher certificate

Convenzione CITES **CITES Convention**

Protocollo di Nagoya Nagoya Protocol

Based on information concerning the raw materials, production process and equipment used they are not likely to be present.

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Based on information concerning the raw materials, production process and equipment used radioactive material is not expected to be present and no irradiation has been used.

The product doesn't contain any nanomaterials according to the new European Cosmetic 1223/2009/EC Regulation and anv nanotechnology is used to produce it

Based on information concerning the raw materials, production process and equipment used polycyclic aromatic hydrocarbons are not likely to be present.

Based on information concerning the raw materials, production process and equipment used they are not likely to be present.

Based on information concerning the raw materials, production process and equipment used they are not likely to be present.

The product is of cosmetic grade and it can be used in cosmetic products.

It is according Regulation 1223/2009, its annexes and its further amendments. We are EFfCI GMP certified.

Yes

Not applicable, cultivated vegetable raw materials

Not applicable

8.1	Data di Retest Retest date	The product, if well preserved and in its original containers, maintains its appearance and characteristics for at least one year from delivery date. After this time, product can be used but it must be rechecked (pH and aspect).
8.2	Stocaggio Storage recommendation	Store at room temperature (15°C-25°C). Protect from frost. At low temperature and on prolonged storage turbidity and stratification phenomena can appear. Indirect warming with stirring will restore the product to its former appearance. This doesn't affect the quality of the product. The viscosity of the product can increase if the storage is higher than 20°C for a long time. Due to its high salt content it can have a corrosive effect during storage in stainless steel tanks. Always homogenize before using.

Data / Date 13/10/2020

Queste informazioni si riferiscono solo al prodotto sopramenzionato e non possono essere considerate valide per altri prodotti o in altri processi produttivi. Le informazioni sono corrette e complete secondo le nostre attuali conoscenze e sono date in buona fede ma senza garanzia. E' responsabilità dell'utilizzatore l'assicurarsi che le informazioni siano appropriate e complete per lo specifico uso del prodotto.

This Information refers only to the above mentioned product and does not need to be valid if used with other product(s) or in any process. The information is to our best present knowledge correct and complete and is given in good faith but without warranty. It remains the user's own responsibility to make sure that the information is appropriate and complete for his specific use of this product.