

Product Documentation

Akoline GCTM

Version
Date 2024 01 10



To whom it may concern

Dear valued customer:

The purpose of this document is to provide you with the information required to evaluate the safety of this product to fulfil the legal requirements. The second purpose of the document is to provide you with all information required during the coding process. AAK has gathered the questions received throughout the years and collected the answers within this document. The document is strictly addressing the cosmetic and personal care applications, thus having no intention to cover, pharmaceutical, food or other applications. As the regulatory requirements increases on the answers given as well as the number of questionnaires increases, AAK has chosen to focus on quality and to give you an answer within a reasonable time. This document represents the answer to your questionnaire. AAK has tried to be as complete and accurate as possible in providing the information and feels comfortable it covers the needs for you. In the case AAK does not possess data or information for a particular subject it is stated in the document.

Head of Development AAK-PC

Staffan Norberg



1.			Contents IDENTIFICATION	8
••	1.1		Identification	· ·
2.			CHEMICAL AND PHYSICAL DATA	9
	2.1		Specifications	·
	2.2		Typical values	
	2.3		Certificate of Analysis	
	2.4		Auxiliary chemical and physical data	
3.			RAW MATERIAL	10
	3.1		Biological data	
	3.2		Composition breakdown	
4.			PRODUCTION	11
	4.1		Production data	
5.			BY-PRODUCTS AND OTHER IMPURITIES	12
	5.1		AAK Contaminant standard	
	5.2		Other impurities specific substanses	
	5.3		Impurities general statements	
		5.3.1	Allergenes	
		5.3.2	Proteins	
		5.3.3	VOC Volatile Organic Compounds	
		5.3.4	Sulphonates	
		5.3.5	Parabens	
		5.3.6	Phthalates	
		5.3.7	Silicones	
6.			STABILITY DATA	13
	6.1		Stability Data	
7.			HUMAN HEALTH HAZARD ASSESSMENT	14



7.1		General read-across consideration and justification
7.2		Acute toxicity
	7.2.1	Acute oral toxicity
	7.2.2	Acute inhalation toxicity
	7.2.3	Acute dermal toxicity
	7.2.4	Acute toxicity by other exposure routes
	7.2.5	Summary and discussion of acute toxicity
7.3		Irritation & corrosivity
	7.3.1	Skin irritation and corrosivity
	7.3.2	Eye & mucous membrane irritation and corrosivity
	7.3.3	Summary and discussion on irritation and corrosivity
7.4		Skin sensitization
	7.4.1	Summary and discussion of sensitisation
7.5		Repeated dose, sub-chronic and chronic toxicity
	7.5.1	Oral administration
	7.5.2	Inhalation studies
	7.5.3	Dermal administration
	7.5.4	Other routes of administration
	7.5.5	Human information
7.6		Reproduction toxicity
	7.6.1	Non-human information
	7.6.2.	Human information
	7.6.3	Developmental toxicity/teratogenicity
	7.6.4	Summary and discussion of reproductive toxicity
7.7		Mutagenicity/genotoxicity
	7.7.1	In vitro data
	7.7.2	In vivo data
	7.7.3	Human information



		7.7.4	Summary and discussion of mutagenicity	
	7.8		Carcinogenicity	
		7.8.1	Non-human information	
		7.8.2	Human information	
		7.8.3	Summary and discussion of carcinogenicity	
	7.9		Toxicokinetics (absorption, metabolism, distribution and elimination (ADME))	
		7.9.1	Oral administration	
		7.9.2	Dermal administration	
		7.9.3	Inhalation route	
	7.10		Photoinduced toxicity	
		7.10.1	Phototoxicity: photoirritation / photosensitisation	
		7.10.2	Phototoxicity: photomutagenicity / photoclastogenicity	
		7.10.3	Other relevant human studies (clinical)	
	7.11		Special investigations	
	7.12		Summary and NOAEL statement	
8.			ECOLOGICAL DATA	37
	8.1		Degradability	
	8.2		Accumulation	
	8.3		Aquatic toxicity	
9.			REGULATORY	41
	9.1		EU	
		9.1.1	EU Cosmetic Regulation EC 1223/2009	
		9.1.2	EU Cosmetic Regulation EC 1223/2009, Annex II and III	
		9.1.3	EU REACH 1907/2006	
		9.1.4	EU SVHC (Substance of Very High Concern)	
		9.1.5	Other	
	9.2		USA	
		9.1.3 9.1.4	EU REACH 1907/2006 EU SVHC (Substance of Very High Concern)	
	9.2	9.2.1	USA US (California) Proposition 65	



		9.2.2	China – NMPA	
		9.2.3	UK REACH	
		9.2.4	Turkey – KKDIK	
		9.2.5	Australia – TGA	
		9.2.6	Other	
	9.3		Other non-Country specific regulatory issues	
		9.3.1	Animal testing	
		9.3.2	Nano particlesTurkey	
		9.3.3	Nagoya Protocol / Biodiversity and Access Benefit Sharing regulation	
		9.3.4	CITES	
		9.3.5	CMR	
		9.3.6	Other	
	9.4		Inventory lists	
10			General statements and standards	43
10	10.1		General statements and standards Official standards	43
10	10.1 10.2			43
10		10.2.1	Official standards	43
10		10.2.1 10.2.2	Official standards Private standards	43
10			Official standards Private standards Ecocert, Cosmos or Natrue	43
10		10.2.2	Official standards Private standards Ecocert, Cosmos or Natrue Vegan and Vegetariam claim	43
10	10.2	10.2.2	Official standards Private standards Ecocert, Cosmos or Natrue Vegan and Vegetariam claim Other	43
10	10.2	10.2.2 10.2.3	Official standards Private standards Ecocert, Cosmos or Natrue Vegan and Vegetariam claim Other Other Statements	43
10	10.2	10.2.2 10.2.3 10.3.1	Official standards Private standards Ecocert, Cosmos or Natrue Vegan and Vegetariam claim Other Other Statements BSE/TSE statements	43
10	10.2	10.2.2 10.2.3 10.3.1 10.3.2	Official standards Private standards Ecocert, Cosmos or Natrue Vegan and Vegetariam claim Other Other Statements BSE/TSE statements GMO statement	43
	10.2	10.2.2 10.2.3 10.3.1 10.3.2	Official standards Private standards Ecocert, Cosmos or Natrue Vegan and Vegetariam claim Other Other Statements BSE/TSE statements GMO statement Other	



	11.3		ISO 9001	
	11.4		EFfCI GMP	
	11.5		Food Safety/ FSSC 22000	
	11.6		Other	
12.			PATENTS	45
	12.1		Patents	
13.			TRANSPORTS AND HANDLING	46
	13.1		Transports	
	13.2		storage unopen package	
	13.3		Handling of product for use	
		13.3.1	Use of full package	
		13.3.2	Use of full package for partly use	
14.			REFERENCES	47
	14.1		References	
15.			DISCLAIMER	48
	15.1		Disclaimer	
16.			APPENDIX	
		C0043	Certificate of Analasys	49
		T0005	Process flowchart	50



1.1 Identification

Producer: AAK Sweden AB, Västra kajen SE-374 82 Karlshamn, Sweden

Tradename: Akoline GC™

Art. No: 8571 Country of Origin EU

This product is used globally. As the product may fit in the definition of several CAS numbers, AAK give examples of alternative CAS number to be used for instance in inventory lists search.

	INCI	CAS Number	EC number
EU /AAK first choice	Hydrogenated Vegetable Glycerides Citrate	97593-31-2	307-334-9
US	Hydrogenated Vegetable Glycerides Citrate	97593-31-2	307-334-9
China*	氢化植物油甘油酯类柠檬酸酯	97593-31-2	
Alternative INCI			

^{*)} For NMPA information see section 9.2.2 China – NMPA

Margrét Viborg

Global Regulatory Affairs Manager



2.1 Specifications

For specification see Product Data Sheet (PDS)

Download latest version at www.aakpersonalcare.com/

2.2 Typical values

For typical values see Product Data Sheet (PDS)

Download latest version at www.aakpersonalcare.com/

2.3 Certificate of Analysis

For example of COA, see Appendix.

2.4 Auxiliary chemical and physical data

Molecular weight ~880 g/mol



3.1 Biological data

Botanical origin

INCI	Botanical origin	*)Geographical origin	Part used	Content %	Wild grown or cultivated
Hydrogenated Vegetable Glycerides Citrate	Elaeis Guineensis	Malaysia or Indonesia	Fruit flesh	100	Cultivated

^{*)}Geographical origin may change

3.2 Composition breakdown

INCI name (EU)	CAS	EINECS	Average Content %	Function
Hydrogenated Vegetable Glycerides Citrate	97593-31- 2	307-334-9	100	Emulsifier

Palm content:			
⊠Containing palm			
□RSPO SG:			
⊠RSPO MB: CU-RSPO SCC-817671			
□Do not contain Palm			



Margrét Viborg Global Regulatory Affairs Manager Personal Care, AAK Sweden AB



4.1 Production data

The following operations are used in the processing of this ingredient

Process		Comment
Mechanical extraction		
Solvent extraction		
Refining		
Deodorising		
Hydrogenation		
Interesterification		
Esterification	Х	
Winterisation		
Solvent Fractionation		
Dry Fractionation		
Ethoxylation		
Molecular distillation		
Other processing	Χ	See attached flowchart



5. BY-PRODUCTS AND OTHER IMPURITIES

5.1 AAK Contaminant standard

Not available, for more information please contact AAK.

5.2 Other Impurities specific substances

Download latest version of "AAK personal Care position on impurities" at aakpersonalcare.com

5.3 Impurities AAK Cosmetic Products

5.3.1 Allergens

Download "General statements AAK Cosmetic Ingredients" at aakpersonalcare.com

5.3.2 Proteins

Download "General statements AAK Cosmetic Ingredients" at aakpersonalcare.com

5.3.3 VOC – Volatile Organic Compounds

Download "General statements AAK Cosmetic Ingredients" at aakpersonalcare.com

5.3.4 Sulphonates

Download "General statements AAK Cosmetic Ingredients" at aakpersonalcare.com

5.3.5 Parabens

Download "General statements AAK Cosmetic Ingredients" at aakpersonalcare.com

5.3.6 Phthalates

Download "General statements AAK Cosmetic Ingredients" at aakpersonalcare.com

5.3.7 Silicones

Download "General statements AAK Cosmetic Ingredients" at aakpersonalcare.com



6.1 Stability data

OSI Value @ 120C No data

Storage @ 20C

Peroxide value 12 month: No data 24 month: No data

Storage @ 40C

Peroxide value 12 month: No data 24 month: No data



7 Human Health and Environmental Hazard Assessment

Akoline GC

7.01 General read-across consideration and justification

Test name:

CIR safety report

Method and laboratory:

Safety assessment and review of various glyceryl ester based cosmetic ingredients

Test material:

Glyceryl stearate citrate and other similar substances

Results:

The panel concludes that existing and available data <u>do not</u> support the safety of glyceryl diesters and their derivatives for use in cosmetic products, mainly due to suspected tumor promoting effect of 1,2-diacylglycerol.

Comments:

See also Amended CIR safety evaluation (S-323).

Read across

Statement

Reference ID:

S-322 Safety assessment of glyceryl dilaurate, glyceryl diarachidate... Final Report of the Cosmetic Ingredient Review Expert Panel, 2002 (available on-line at https://www.cir-safety.org/)

Test name:

Amended CIR safety report

Method and laboratory:

Safety assessment and review of various glyceryl ester based cosmetic ingredients

Test material:

Glyceryl stearate citrate and other similar substances

Results:

The panel concluded that glyceryl dilaurate, glyceryl diarachidate, glyceryl stearate citrate and other substances covered in the amended review are safe for use in cosmetics, provided that the level of 1,2-diesters of glycerol is not high enough to induce epidermal hyperplasia.

Comments:

The permitted level of 1,2-diesters is not explicitly stated. In view of the rapid hydrolysis by lipases and the higher stability of 1,3-diesters versus 1,2-diesters it is unlikely that the level of 1,2-diesters in actual applications will be high enough to be of concern.

Read across

Statement

Reference ID:

S-323 Amended final report on the safety assessment of glyceryl dilaurate, glyceryl diarachidate..., Johnson, W, Int J Toxicol, 26 (Suppl 3), 1-30, (2007)



Test name:

EFSA Safety assessment of several polar derivatives of mono- and diglycerides

Method and laboratory:

Opinion and statement from EFSA on the safety of use of citric acid esters of mono- and diglycerides as food emulsifiers. The report also comprises a summary of existing toxicological studies on other derivatives of mono- and diglycerides (E472a-f).

Test material:

Citric acid esters of mono- and diglycerides (CITREM, E472c)

Results:

There are very few relevant studies for E472c in the EFSA safety assessment. However, due to the hydrolysis of the citric acid ester to citric acid and mono- and diglycerides (which are further metabolized to fatty acids and glycerol), it is concluded that E472c is safe for use as a food additive and no limit on intake (ADI) is necessary.

Read across

Statement

Reference ID:

S-338 Re-evaluation of acetic acid, lactic acid, citric acid, tartaric acid, mono- and diacetyltartaric acid, mixed acetic and tartaric acid esters of mono- and diglycerides of fatty acids (E 472a-f) as food additives, EFSA Journal 18(3):6032, 1-66

Test name:

REACH registration dossier

Method and laboratory:

REACH registration dossier with toxicological and environmental data

Test material:

Glycerides, C16-18 and C18-unsatd., mono- and di-, citrates, (CAS 91052-16-3)

Results:

See separate entries for details

Read across

Statement

Reference ID:

S-339 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono- and dicitrates', CAS number 91052-16-3

https://echa.europa.eu/registration-dossier/-/registered-dossier/18167

Test name:

REACH registration dossier

Method and laboratory:

REACH registration dossier with toxicological and environmental data

Test material:

Glycerides, C16-18 and C18-unsatd., mono-, di- and tri-, citrates, potassium salts, (CAS 91744-23-9)

Read across

Statement



Reference ID:

S-340 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono-, di- and tri-, citrates, potassium salts', CAS number 91744-23-9

https://echa.europa.eu/registration-dossier/-/registered-dossier/24811

Test name:

REACH registration dossier

Method and laboratory:

REACH registration dossier with toxicological and environmental data

Test material:

Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates

Read across

Statement

Reference ID:

S-341 ECHA, REACH registration 'Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates'

https://echa.europa.eu/registration-dossier/-/registered-dossier/23747

Test name:

REACH registration dossier

Method and laboratory:

REACH registration dossier with toxicological and environmental data

Test material:

Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, potassium salts, (CAS 91744-38-6)

Read across

Statement

Reference ID:

S-342 ECHA, REACH registration 'Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, potassium salts', CAS 91744-38-6

https://echa.europa.eu/registration-dossier/-/registered-dossier/12022

Test name:

REACH registration dossier

Method and laboratory:

REACH registration dossier with toxicological and environmental data

Test material:

Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, sodium salts, (CAS 91744-39-7)

Read across

Statement

Reference ID:

S-343 ECHA, REACH registration 'Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, sodium salts', CAS 91744-39-7

https://echa.europa.eu/registration-dossier/-/registered-dossier/12131



Citric acid esters of mono- and diglycerides belong to a group of ingredients that share the mono- and diglyceride structure as a common denominator. Glycerol is esterified to one or two straight chain saturated or unsaturated fatty acids, normally a combination of palmitic and stearic acid

The free hydroxyl groups on the mono- and diglycerides are further reacted with an acid, ranging from acetic to tartaric acid. These ingredients are usually used as food additives, as emulsifying agents or conditioning agents for prepared food. One big use area is powders for infant nutrition. This class of food additives is known as E472a-f, where (a) is the acetic acid derivative, (b) is lactic acid esters, (c) is citric acid esters and (d)-(f) are variations containing tartaric acid.

The chemistry of this group of compounds is rather similar from a toxicological point of view. The esters are usually rapidly hydrolyzed in the stomach and intestines by lipases to the corresponding acids and glycerol. The citric acid esters of mono- and diglycerides would typically release citric acid, palmitic acid and stearic acid as well as glycerol, which are all endogenous substances in normal metabolism. They are generally regarded as non-toxic functional ingredients for food applications and very few new studies have been published in the last 20 years.

The nomenclature, INCI names and CAS numbers for the citric acid esters of mono- and diglycerides vary depending on the producer of the material (See 7.11 for a summary). For saturated mono- and diglycerides both "hydrogenated vegetable glycerides citrate" and "glyceryl stearate citrate" are used. In both cases the names should be interpreted as "mixed mono- and diglycerides, comprising palmitic and stearic acid in different proportions, esterified with citric acid". From a toxicological point of view, these variations are insignificant.

The citric acid esters of mono- and diglycerides can be partially or completely neutralized by a weak alkali resulting in sodium or potassium salts. These sodium or potassium salts are easier to disperse in water than the corresponding acid form and are often preferred due to the higher functionality in the application. From a toxicological point of view the neutralised and the acid forms can be seen as equal.

The two CIR safety reports from 2002 and 2007 are mainly dealing with underivatized monoand diglycerides of fatty acids and are of limited use for the evaluation of the citric acid esters, especially since the data reported are old and without proper references to original articles. Only data where enough details about test methodology are listed are presented in this safety report.

Several entries of citric acid esters of mono- and diglycerides are found in REACH registrations. The toxicological data and environmental toxicity data are heavily read-across and only one registration is extensively extracted in this safety review.



7.02 Acute toxicity7.02.1 Acute oral toxicity

Test name:

Acute oral toxicity

Method and laboratory:

OECD Guideline 423, OECD Guideline 401

Test material:

Glycerides, C16-18 & C18-unsatd., mono- and di-, acetates, (CAS 91052-13-0) Glyceryl citrate/lactate/linoleate/oleate

Results:

LD50>2000 mg/kg bw

Read across

Read across from structurally similar substances

Reference ID:

S-339 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono- and dicitrates', CAS number 91052-16-3

https://echa.europa.eu/registration-dossier/-/registered-dossier/18167

7.02.2 Acute inhalation toxicity

No actual tests have been carried out and literature data has not been found for this chapter.

7.02.3 Acute dermal toxicity

Test name:

Acute dermal toxicity

Method and laboratory:

OECD Guideline 402

Test material:

Glycerides, C16-18 & C18-unsatd., mono- and di-, acetates, (CAS 91052-13-0) Glyceryl citrate/lactate/linoleate/oleate

Results:

LD50>2000 mg/kg bw

Read across

Read across from structurally similar substances

Reference ID:

S-339 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono- and dicitrates', CAS number 91052-16-3

https://echa.europa.eu/registration-dossier/-/registered-dossier/18167

7.02.4 Acute toxicity by other exposure routes

No actual tests have been carried out and literature data has not been found for this chapter.



7.02.5 Summary and discussion of acute toxicity

No fully verified data for the acute oral, dermal or inhalation toxicity has been identified for Akoline GC/LC. The CIR report states the LD50 (Oral) and LD50 (dermal) to be >2 g/kg bw/day but no published data are shown. It is stated in the EFSA report that although no data exist for the acute oral toxicity, absence of short term and sub-chronic toxicity indicate that acute toxicity would not be a concern. The various REACH registrations for citric acid esters of mono- and diglycerides report read-across studies, and values of LD50(Oral) / LD50(dermal) in excess of 2000 mg/kg bw are reported.

It is concluded that acute toxicity is not a health hazard or concern for this substance group.



7.03 Irritation & corrosivity

7.03.1 Skin irritation and corrosivity

Test name:

Summary of several skin irritation studies in REACH registration

Method and laboratory:

OECD Guideline 404. Rabbits, semi-occlusive patch, duration of exposure 4h, observation up to 7-14 days.

Test material:

Glyceryl citrate/lactate/linoleate/oleate

Results:

Not irritating

Read across

Read across from structurally similar substances

Reference ID:

S-339 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono- and dicitrates', CAS number 91052-16-3

https://echa.europa.eu/registration-dossier/-/registered-dossier/18167

Test name:

Skin irritation (24h human closed patch test)

Method and laboratory:

No guideline followed. N=40. Dilution 1:10 in water. Application of closed patch for 24 h, evaluation at 24, 48 and 72 h after removal of patch.

Test material:

Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, potassium salts, (CAS 91744-38-6)

Results:

Not irritating

Read across

Read across from structurally similar substances

Reference ID:

S-342 ECHA, REACH registration 'Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, potassium salts', CAS 91744-38-6

https://echa.europa.eu/registration-dossier/-/registered-dossier/12022

7.03.2 Eye & mucous membrane irritation and corrosivity

Test name:

Summary of eye irritation in REACH registration

Method and laboratory:

OECD Guideline 405. Rabbits, exposure 24 hours, observations up to 72 hours.

Test material:

Glyceryl citrate/lactate/linoleate/oleate



Results:

Not irritating

Read across

Read across from structurally similar substances

Reference ID:

S-339 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono- and dicitrates', CAS number 91052-16-3

https://echa.europa.eu/registration-dossier/-/registered-dossier/18167

7.03.3 Summary and discussion on irritation and corrosivity

No actual skin irritation or eye irritation studies have been performed on Akoline GC/LC. The CIR safety reports mention old unpublished studies from Huls America Inc, with slight indication of skin irritation and no eye irritation but study details are not available. From the read-across data for structurally similar substances in the REACH registrations, it can be concluded that skin or eye irritation is not likely to occur.

It is therefor concluded that citric acid esters of mono- and diglycerides are not irritating to the skin or the eye and mucous membranes.

In the absence of skin/eye irritation, it is concluded that citric acid esters of mono- and diglycerides are not likely to cause skin or eye corrosion.



7.04 Skin sensitization

Test name:

Summary of skin sensitization data in REACH registration

Method and laboratory:

Several in vivo tests (OECD 406, Magnusson Kligman Maximization Test) are reported. The test substances are typically diluted in mineral oil for induction tests and petrolatum for challenge tests.

Test material:

Glycerides, C16-18 and C18-hydroxy mono- and di-, (CAS 91845-19-1) Glycerides C16-18 mono-, di-and tri-, hydrogenated, citrates, potassium salts, (CAS 91744-38-5)

Results:

Not sensitizing

Read across

Read across from structurally similar substances

Reference ID:

S-339 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono- and dicitrates'. CAS number 91052-16-3

https://echa.europa.eu/registration-dossier/-/registered-dossier/18167

Test name:

In vivo skin sensitization test

Method and laboratory:

Magnusson-Kligman guinea pig sensitization test. N=10.Challenge by intradermal injection of 5% glyceryl stearate citrate in paraffin oil, challenge after 14 days by topical application of 25% glyceryl stearate citrate in petrolatum.

Laboratory: Safepharm Laboratories, 1981.

Test material:

Glyceryl stearate citrate

Results:

Not sensitizing

Read across

Read across from structurally similar substances

Reference ID:

S-323 Amended final report on the safety assessment of glyceryl dilaurate, glyceryl diarachidate..., Johnson, W, Int J Toxicol, 26 (Suppl 3), 1-30, (2007)

Test name:

Skin sensitization

Method and laboratory:

OECD 406 (Buehler/Magnusson-Kligman). Induction and challenge 2, 25, 50 and 100% in corn oil. Challenge with undiluted test substance on day 28.



Test material:

Glycerides, C16-18 and C18-unsatd., mono-, di- and tri-, citrates, potassium salts, (CAS 91744-23-9)

Results:

Not sensitizing

Read across

Read across from structurally similar substances

Reference ID:

S-340 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono-, di- and tri-, citrates, potassium salts', CAS number 91744-23-9

https://echa.europa.eu/registration-dossier/-/registered-dossier/24811

7.04.1 Summary and discussion of sensitization

According to read-across studies, presented in the CIR safety report and in REACH registration dossiers, citric acid esters of mono- and diglycerides are not sensitizing.



7.05 Repeated dose, sub-chronic and chronic toxicity

7.05.1 Oral administration

Test name:

Summary of repeated dose toxicity studies

Method and laboratory:

6 separate studies are presented in the document. Only one study was performed with citric acid esters of mono- and diglycerides (1966). This 10 day study comprises two treatment levels (23100 and 37500 mg/kg bw/day), with lard as reference substance. Mortalities, organ weights, gross pathologies and histopathologies were investigated. The NOAEL based on this study was set to >= 37500 mg/kg bw/day.

Test material:

Castor oil, (CAS 8001-79-4)

Glycerides, C8-18 and C18-unsatd., mono- and di-, acetates, (CAS 92052-13-0)

Glycerides, C16-18 and C18-hydroxy, mono- and di-, (CAS 91845-19-1)

Glycerides, C16-18 and C18-unsatd., mono- and di-, citrates (CAS 91052-16-3)

Results:

It is concluded on the basis of the studies on the read-across substances and the citric acid esters, that there are no observed adverse effects and no repeated dose toxicity hazards are identified.

Comments:

Also referenced in the EFSA evaluation (S-338)

Read across

Read across from structurally similar substances

Reference ID:

S-339 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono- and dicitrates', CAS number 91052-16-3

https://echa.europa.eu/registration-dossier/-/registered-dossier/18167

7.05.2 Inhalation studies

No actual tests have been carried out and literature data has not been found for this chapter.

7.05.3 Dermal administration

No actual tests have been carried out and literature data has not been found for this chapter.

7.05.4 Other routes of administration

No actual tests have been carried out and literature data has not been found for this chapter.

7.05.5 Human studies

No actual tests have been carried out and literature data has not been found for this chapter.



7.05.6 Summary and discussion

The EFSA evaluation of citric acid esters of mono- and diglycerides conclude that the short term and sub-chronic toxicity is very low and no ADI (Acceptable Daily Intake) value is required in food applications. Only one actual study from 1966 on citric acid esters of mono- and diglycerides, stating a NOAEL (oral, systemic) to be as high as 37500 mg/kg bw/day. The available REACH registrations refer to read-across from similar substances.

It is concluded that the citric acid esters of mono- and diglycerides have very low short term and sub-chronic toxicity.



7.06 Reproduction toxicity

7.06.1 Non-human studies

Test name:

Summary of reproductive and developmental toxicity

Method and laboratory:

4 different studies on the reproductive toxicity and 8 studies on developmental toxicity are presented in the summary. No data on citric acid esters of mono- and diglycerides were found so read-across from structurally similar substances is used as the base for the evaluation.

Test material:

Glycerides, castor-oil, mono-, hydrogenated, acetates, (CAS 736150-63-3) Glycerides, C8-18 and C18-unsatd., mono- and di-, acetates, (CAS 91052-13-0)

Results:

No adverse effects on the reproductive health or fertility were observed. No adverse effects on the developmental toxicity were observed.

Read across

Read across from structurally similar substances

Reference ID:

S-339 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono- and dicitrates', CAS number 91052-16-3

https://echa.europa.eu/registration-dossier/-/registered-dossier/18167

7.06.2 Human studies

No actual tests have been carried out and literature data has not been found for this chapter.

7.06.3 Developmental toxicity/teratogenicity

7.06.3.1 Non-human studies

No actual tests have been carried out and literature data has not been found for this chapter.

7.06.3.2 Human studies

No actual tests have been carried out and literature data has not been found for this chapter.

7.06.4 Summary and discussion of reproductive toxicity

No published studies on citric acid esters of mono- and diglycerides regarding reproductive and developmental toxicity have been identified. The read-across studies presented in REACH registration documents and in the EFSA evaluation state that no treatment related adverse effects on reproduction or off-spring development have been identified, up to the highest tested levels. As the highest tested levels vary from about 1000 mg/kg bw/day to about 8600 mg/kg bw/day in the different studies, a NOAEL is difficult to assign to the citric acid esters of mono- and diglycerides. The relevance of the the read-across substances also need to be considered.



The studies presented in the EFSA evaluation and in the REACH registrations conclude that reproductive and developmental toxicity is low for citric acid esters of mono- and diglycerides, based on read-across data from related substance classes. Acetic acid esters of castor oil glycerides showed NOAEL values of 1159-2200 mg/kg bw/day (M/F) for fertility and 1342-2260 mg/kg bw/day (generation F1) for fetal development. Acetic acid esters of C8-18 & C18-unsaturated mono/diglycerides showed a NOAEL of 1000 mg/kg bw/day for both fertility and developmental toxicity. In both cases the NOAEL represented the highest tested dose.

It is concluded that citric acid esters of mono- and diglycerides can be considered non-toxic for reproduction and development and a conservative estimation of the NOAEL can be set to 2200 mg/kg bw/day.



7.07 Mutagenicity/genotoxicity

7.07.1 In vitro data

Test name:

Summary of mutagenicity and genotoxicity tests

Method and laboratory:

9 different studies are presented for different citric acid esters of mono- and diglycerides and from structurally similar substances. The tests comprise in vitro mammalian chromosome aberration tests, in vitro gene mutation study in bacteria (Ames' test, with and without metabolic activation) and in vitro gene mutation tests in mammalian cells.

Test material:

Glycerides, C8-18 and C18-unsatd., mono- and di-, acetates, (CAS 92052-13-0) Glyceryl citrate/lactate/linoleate/oleate Glycerides, C16-18 mono- di- and tri-, hydrogenated, citrates, potassium salts, (CAS 91744-38-6)

Results:

All the tested substances are found to be non-mutagenic.

Comments:

See reference for additional tested substances.

Read across

Read across from structurally similar substances

Reference ID:

S-339 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono- and dicitrates', CAS number 91052-16-3

https://echa.europa.eu/registration-dossier/-/registered-dossier/18167

Test name:

In vitro genotoxicity

Method and laboratory:

OECD 476 (In vitro Mammalian Cell Gene Mutation Test). Chinese hamster lung fibroblasts, with and without metabolic activation. Dosage 4.7-150 microgram/l in THF.

Test material:

Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates

Results:

Not genotoxic

Read across

Read across from structurally similar substances

Reference ID:

S-341 ECHA, REACH registration 'Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates'

https://echa.europa.eu/registration-dossier/-/registered-dossier/23747



7.07.2 In vivo data

No actual tests have been carried out and literature data has not been found for this chapter.

7.07.3 Human studies

No actual tests have been carried out and literature data has not been found for this chapter.

7.07.4 Summary and discussion of mutagenicity

Based on in vitro studies on gene mutation tests in bacteria and mammalian cells as well as mammalian chromosome aberration tests on substances that are structurally relevant, it is concluded that citric acid esters of mono- and diglycerides do not present any concern for mutagenicity and genotoxicity.



7.08 Carcinogenicity

7.08.1 Non-human studies

No actual tests have been carried out and literature data has not been found for this chapter.

7.08.2 Human studies

No actual tests have been carried out and literature data has not been found for this chapter.

7.08.3 Summary and discussion of carcinogenicity

No published studies have been found for carcinogenic effects of citric acid esters of monoand diglycerides or relevant structurally similar substances.

The CIR safety assessments from 2002 and 2007 (amended) refer to potential risks involved in diglycerides causing or promoting epidermal hyperplasia (abnormal increase in number of cells). The effect is said to be linked to 1,2-diglycerides, especially with short chain fatty acids (C10, C12) or mixed saturated/unsaturated fatty acids. The citric acid esters of mono- and diglycerides that are reviewed in this safety assessment are based on C16 and C18 saturated fatty acids and do not contain appreciable amounts of unreacted 1,2-diglycerides of either short chain or mixed saturated/unsaturated fatty acids.

Due to the rapid metabolism in the skin by microbial or endogenous lipases, the citric acid esters of mono- and diglycerides are hydrolyzed into free fatty acids (palmitic & stearic acid), citric acid and glycerol. All of these substances are part of a normal mammalian metabolism and do not constitute known risks for carcinogenicity.

It is concluded that citric acid esters of mono- and diglycerides are unlikely to be a health hazard from a carcinogenicity point of view.



7.09 Toxicokinetics: absorption, metabolism, distribution and elimination (ADME)

7.09.1 Oral administration

Test name:

Toxicokinetics (ADME)

Method and laboratory:

Summary of toxicokinetic data (absorption, distribution, metabolism and excretion) for citric acid esters of mono- and diglycerides and other structurally similar glycerides.

Test material:

None

Results:

No actual data on the ADME have been identified. It is postulated that citric acid esters of mono- and diglycerides undergo the same type of absorption and metabolism as normal triglycerides, ie hydrolysis aided by lipases into free fatty acids, citric acid and glycerol, which are further metabolized and excreted.

Comments:

Uptake via the dermal route is indicated to be low, on grounds of the molecular weight and polarity of the substance.

Uptake via the inhalation route can occur if the substance is present in aerosol particles with a diameter <15 micrometers.

Read across

Read across Generic discussion of ADME of glycerides, including citrates and

other similar derivatives of mono- and diglycerides

Reference ID:

S-339 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono- and dicitrates', CAS number 91052-16-3

https://echa.europa.eu/registration-dossier/-/registered-dossier/18167

Test name:

In vitro digestion study

Method and laboratory:

In vitro investigation on the metabolism of citric acid esters of mono- and diglycerides in simulated infant food application. The citric acid esters are studied both alone and in a simulated infant food emulsion.

Test material:

Citric acid ester of mono- and diglycerides, (Grindsted CITREM LR10, Danisco A/S)

Results:

The citric acid ester is rapidly hydrolyzed to free fatty acids and a glyceryl citrate. The water-soluble glyceryl citrate can be persistent in some conditions in the in vitro study.

Read across

Read across from structurally similar substances

Reference ID:

S-325 In vitro digestion of citric acid esters of mono- and diglycerides (CITREM) and CITREM-containing infant formula/emulsions, Amara, S. et al, Food&Function, 5(7), 1409-1421 (2014)



7.09.2 Dermal administration

No actual tests have been carried out and literature data has not been found for this chapter.

7.09.3 Inhalation route

No actual tests have been carried out and literature data has not been found for this chapter.



7.10 Photoinduced toxicity

No published studies on the phototoxicity, photoirritation, photosensitization or other photoinduced effects for citric acid esters of mono- and diglycerides or relevant read-across substances have been found.

The chemical structure of citric acid esters of mono- and diglycerides and the absence of photo-induced toxicity for the constituents makes it unlikely that any health hazards linked to photo-toxicity would occur at normal use conditions.

7.10.1 Phototoxicity: photoirritation / photosensitisation

No actual tests have been carried out and literature data has not been found for this chapter.

7.10.2 Phototoxicity: photomutagenicity / photoclastogenicity

No actual tests have been carried out and literature data has not been found for this chapter.

7.10.3 Other relevant human studies (clinical)

No actual tests have been carried out and literature data has not been found for this chapter.



7.11 Special investigations

The citric acid esters of mono- and diglycerides are common emulsifiers and food additives, as well as functional ingredients in personal care products. Due to the different origins and backgrounds, the nomenclature is complicated. This section is listing the different names and identification numbers for the citrates covered by this safety report.

INCI name: CAS number(s):

Glyceryl stearate citrate: 55840-13-6, 86418-55-5, 39175-7-9

Hydrogenated palm glycerides citrate: 91744-68-2
Hydrogenated vegetable glycerides citrate: 97593-31-2
Hydrogenated tallow glyceride citrate: 68990-59-0
Hydrogenated tallow glycerides citrate: 91723-33-0
Glyceryl citrate/lactate/linoleate/oleate: No CAS found

REACH nomenclature:

Glycerides, C16-18 and C18-unsatd., mono- and di-, citrates: 91052-16-3 Glycerides, C16-18 and C18-unsatd., mono- and di-, citrates, calcium salts: 91052-18-5 Glycerides, C16-18 and C18-unsatd., mono- and di-, citrates, potassium salts: 91052-20-9 Glycerides, C16-18 and C18-unsatd., mono- and di-, citrates, sodium salts: 91052-22-1 Glycerides, C16-18 and C18-unsatd., mono-, di- and tri-, citrates, potassium salts: 91744-23-9 Glycerides, C16-18 and C18-unsatd., mono-, di- and tri-, citrates, sodium salts: 91744-24-0 Glycerides, C16-18 and C18-unsatd., mono-, dicitrates: 91052-30-1 Glycerides, C16-18 and C18-unsatd., mono-, dicitrates, calcium salts: 91052-32-3 Glycerides, C16-18 and C18-unsatd., mono-, dicitrates, potassium salts: 91052-34-5 Glycerides, C16-18 mono-, di- and tri-, citrates: 91744-36-4 Glycerides, C16-18 mono-, di- and tri-, hydrogenated, citrates: No CAS found Glycerides, C16-18 mono-, di- and tri-, hydrogenated, citrates, potassium salts: 91744-38-6 Glycerides, C16-18 mono-, di- and tri-, hydrogenated, citrates, sodium salts: 91744-39-7



7.12 Summary and NOAEL statement

Based on the data presented in chapter 7.1-7.11 a NOAEL value of 2200 mg/kg bw/day for systemic exposure is given for citric acid esters of mono- and diglycerides.

This value is based on data presented in Chapter 7.06 (Reproduction and developmental toxicity).



OECD 202 (Daphnia magna, 48h, WAF, 100 mg/l) OECD 201 (Pseudokirchneriella subcapitata, 72h, WAF,6.25-100 mg/l)

Test material:

Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, potassium salts, (CAS 91744-38-6)

Results:

NOELR(fish) 100 mg/l NOELR(Daphnia) 100 mg/l WAF NOELR(algae) 100 mg/l WAF

Read across

Read across from structurally similar substances

Reference ID:

S-342 ECHA, REACH registration 'Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, potassium salts', CAS 91744-38-6 https://echa.europa.eu/registration-dossier/-/registered-dossier/12022

8.04 Summary of ecotoxicity

The citric acid esters of mono- and diglycerides are readily biodegraded, do not accumulate in the environment and show low toxicity to aquatic organisms (fish, invertebrates and algae). It is therefor concluded that they pose minimal risk to the environment.



8 Ecological data8.01 Degradability

Test name:

Biodegradability in water

Method and laboratory:

OECD 301F

Test material:

Glycerides, C16-18 and C18-unsatd., mono- and di-, citrates, (CAS 91052-16-3)

Results:

Readily biodegradable

Read across

Read across from structurally similar substances

Reference ID:

S-339 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono- and dicitrates', CAS number 91052-16-3

https://echa.europa.eu/registration-dossier/-/registered-dossier/18167

Test name:

Biodegradability

Method and laboratory:

OECD 301D (Closed bottle test)

Test material:

Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, sodium salts, (CAS 91744-39-7)

Results:

Readily biodegradable

Read across

Read across from structurally similar substances

Reference ID:

S-343 ECHA, REACH registration 'Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, sodium salts', CAS 91744-39-7

https://echa.europa.eu/registration-dossier/-/registered-dossier/12131

Test name:

Biodegradability

Method and laboratory:

OECD 301D

Test material:

Glycerides, C16-18 and C18-unsatd., mono-, di- and tri-, citrates, potassium salts, (CAS 91744-23-9)

Results:

Readily biodegradable

Read across

Read across from structurally similar substances



Reference ID:

S-340 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono-, di- and tri-, citrates, potassium salts', CAS number 91744-23-9

https://echa.europa.eu/registration-dossier/-/registered-dossier/24811

Test name:

Biodegradability

Method and laboratory:

OECD 301D (Closed bottle test)

Test material:

Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, potassium salts, (CAS 91744-38-6)

Results:

Readily biodegradable

Read across

Read across from structurally similar substances

Reference ID:

S-342 ECHA, REACH registration 'Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, potassium salts', CAS 91744-38-6

https://echa.europa.eu/registration-dossier/-/registered-dossier/12022

8.02 Accumulation

Citric acid esters of mono- and diglycerides as well as relevant read-across substances are readily biodegradable. They are generally easily hydrolyzed to free fatty acids, citric acid and glycerol by aquatic and soil microorganisms. The fatty acids, citric acid and glycerol are easily metabolized by aquatic and soil microorganisms. Therefor the risk of environmental accumulation is regarded as minimal.



8.03 Aquatic toxicity

Test name:

Summary of aquatic toxicity

Method and laboratory:

One long-term study on aquatic invertebrates (daphnia), and two short-term studies (fish & algae) are summarized (OECD 203).

Test material:

Glycerides, C16-18, mono-, di- and tri-, hydrogenated, potassium salts, (CAS 91744-38-6)

Results:

NOEL(21d) = 60 mg/l (daphnia) LL50(96h) > 100 mg/l (fish) NOELR(72h) > 100 mg/l (algae)

Comments:

Conclusion: no aquatic toxicity observed up to solubility limit (about 1.3 mg/l)

Read across

Read across from structurally similar substances

Reference ID:

S-339 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono- and dicitrates', CAS number 91052-16-3

https://echa.europa.eu/registration-dossier/-/registered-dossier/18167

Test name:

Summary of aquatic toxicity tests

Method and laboratory:

OECD 203 (Danio rerio, 96h, WAF, 100 mg/l)

OECD 202 (Daphnia magna, 48h, WAF, 100 mg/l)

OECD 201 (Pseudokirchneriella subcapitata, 72h, WAF,6.25-100 mg/l)

Test material:

Glycerides, C16-18 and C18-unsatd., mono-, di- and tri-, citrates, potassium salts, (CAS 91744-23-9)

Results:

NOELR(fish) 100 mg/l

NOELR(Daphnia) 100 mg/l WAF

NOELR(algae) 100 mg/l WAF

Read across

Read across from structurally similar substances

Reference ID:

S-340 ECHA, REACH registration 'Glycerides, C16-18 and C18-unsatd., mono-, di- and tri-, citrates, potassium salts', CAS number 91744-23-9

https://echa.europa.eu/registration-dossier/-/registered-dossier/24811

Test name:

Aquatic toxicity

Method and laboratory:

OECD 203 (Danio rerio, 96h, WAF, 100 mg/l)



OECD 202 (Daphnia magna, 48h, WAF, 100 mg/l) OECD 201 (Pseudokirchneriella subcapitata, 72h, WAF,6.25-100 mg/l)

Test material:

Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, potassium salts, (CAS 91744-38-6)

Results:

NOELR(fish) 100 mg/l NOELR(Daphnia) 100 mg/l WAF NOELR(algae) 100 mg/l WAF

Read across

Read across from structurally similar substances

Reference ID:

S-342 ECHA, REACH registration 'Glycerides, C16-18, mono-, di- and tri-, hydrogenated, citrates, potassium salts', CAS 91744-38-6 https://echa.europa.eu/registration-dossier/-/registered-dossier/12022

8.04 Summary of ecotoxicity

The citric acid esters of mono- and diglycerides are readily biodegraded, do not accumulate in the environment and show low toxicity to aquatic organisms (fish, invertebrates and algae). It is therefor concluded that they pose minimal risk to the environment.



9.1 EU

9.1.1 Statement on EU Cosmetic Regulation EC 1223/2009

Latest statement, download "Statement on EU Cosmetic Regulation" at aakpersonalcare.com

9.1.2 EU Cosmetic Regulation EC 1223/2009, Annex II and III

Latest statement, download "Statement on EU Cosmetic Regulation" at aakpersonalcare.com

9.1.3 EU REACH 1907/2006

Latest statement, download "REACH Statement" at aakpersonalcare.com

9.1.4 EU SVHC (Substance of Very High Concern)

Latest statement, download "General Statement AAK Ingredients" at aakpersonalcare.com

9.2 Other country specific regulations:

9.2.1 US (California) Proposition 65

Latest statement, download "General Statement AAK Ingredients" at aakpersonalcare.com

9.2.2 China - NMPA

Latest statement, download "NMPA Statement" at <u>aakpersonalcare.com</u>

9.2.3 UK REACH

Latest statement, download "UK REACH Statements" at aakpersonalcare.com

9.2.4 Turkey - KKDIK

Latest statement, download "Turkey-KKDIK and SEA Statement" at aakpersonalcare.com

9.2.5 Australia - TGA

Latest statement, download "AAK PC Products and TGA status" at aakpersonalcare.com

9.3 Other non-Country specific regulatory issues

9.3.1 Animal testing

Latest statement, download "General Statement AAK Ingredients" at <u>aakpersonalcare.com</u>

9.3.2 Nano particles

Latest statement, download "General Statement AAK Ingredients" at aakpersonalcare.com

9.3.3 Nagoya Protocol / Biodiversity and Access Benefit Sharing regulation

Latest statement, download "General Statement AAK Ingredients" at aakpersonalcare.com

9.3.4 CITES

Latest statement, download "General Statement AAK Ingredients" at aakpersonalcare.com

9.3.5 CMR

Latest statement, download "General Statement AAK Ingredients" at aakpersonalcare.com



9.4 Inventory lists

Inventory lists relates to substances already existing in a specific market. The inventory list to the chemical legislation of the country or region. INCI labeling is not related to the chemical legislation. The nomenclature may differ between these two types of regulations hence the wording may change.

In the Table below, column 3:

- 1) Listed means:
 - a. The substance name and CAS number described as "AAK first choice name", in section "1.1 Identification" is listed and not prohibited in the inventory list of the country.
- 2) Not listed, however CAS. No XXXXX-XX-X is listed and valid to be used.
 - a. The substance name and CAS number described as "AAK first choice name", section "1.1 identification" is not found but instead the Cas XXXXX-XX-X mentions is listed as well as fits with the chemical description of the product, hence can be used instead.
- 3) No data:
 - a. AAK has not been able to find the substance in the inventory list.

EC (EU)	EC-inventory	Listed
TSCA (U.S.)	Toxic Substances Control Act	No
DSL (Canada)	Domestic Substances List	No but on the ICL list CAS 55840-13-6 is found.
AICS (Australia)	The Australian Inventory of	No data
	Chemical Substances	
IECSC (China)	Inventory of Existing Chemical	No data
	Substances Produced or	
	Imported in China	
IECIC (China)	Inventory of Existing Cosmetic	Listed
	Ingredients in China	
ENCS (Japan)	Combined list of existing and	No data
	notified chemical substances as	
	the Japanese Existing and New	
	Chemical Substances Inventory.	
Japan	Japan Pharmacopoeia	Not found
KECI (South	Korea Existing Chemicals	No data
Korea)	Inventory	
PICCS	Philippine Inventory of Chemicals	No data
(Philippines)	and Chemical Substances	
NZIoC (New	New Zealand Inventory of	No data
Zealand)	Chemicals	
NECI (Taiwan)	National Existing Chemical	CAS 55840-13-6 is found and valid to be used.
	Inventory	
Saudia Arabia	The Saudi Arabian Standards	No data
	Organisation	
Malaysia	Chemicals Information	No data
	Management System	
Mexico	Inventario Nacional de Sustancias	No data
	Químicas	
Turkey		No data



10.1 Official standards

Standard	Conform	Monograph
EUR/Ph	n.a	
USP/NF	n.a	
JP	See inventory list 9.4	

10.2 Private standards

10.2.1 Ecocert, Cosmos or Natrue

Latest statement, download "General Statement AAK Ingredients" at aakpersonalcare.com

10.2.2 Vegan and Vegetariam claim

Latest statement, download "General Statement AAK Ingredients" at aakpersonalcare.com

10.2.3 Other

10.3 Other Statements

10.3.1 BSE/TSE statements:

Not available, please contact AAK for more information.

10.3.2 GMO statement

Not available, please contact AAK for more information.

10.3.3 Other:

No data



11. CERTIFICATES

11.1 Halal

Contact AAK for more information

.

11.2 Kosher

Contact AAK for more information

11.3 ISO 9001

Contact AAK for more information

11.4 EFFCI GMP

No data

11.5 Food Safety/ FSSC 22000

Contact AAK for more information

11.6 Other

No other available



12. PATENTS

12.1 Patents

No data.



TRANSPORTS AND HANDLING - Akoline GC™

13.1 Transports

No data available

13.2 storage unopen package

Storage to fulfill shelf life:

Store in temperature below 20C or lower. Dark, dry and odour free condition in unopen packaging's. See Product data sheet for more information.

Retest of batch:

Retest for prolonged shelf life is only possible after agreement with sales responsible.

13.3 Handling of product for use

13.3.1 Use of full package

Recommended melting temperature.

Bag is possible to use amount needed directly from the bag. Do not melt material directly in the bag remove and melt in a vessel.

During processing need to be heated to 65C to remove crystal memory.

13.3.2 Use of full package for partly use

Bag is possible to use partly directly from the bag. Do not melt material directly in the bag remove and melt in a vessel.

Reseal packaging and store in 20C or below

Note:

AAK's shelf life for ingredients that are unopened and stored according to the instructions given in the product information sheet. This guarantee is invalidated once the packaging is opened and the ingredients reheated. It is the user's responsibility to validate that a reheated material fulfills shelf life requirements in a formulation. See Product Data Sheet.



14. REFERENCES

14.1 References

No data



15. DISCLAIMER

15.1 Disclaimer

This document, or any answers or information provided herein by AAK, does not constitute a legally binding document of AAK. While the description designs, data and information contained herein are presented in good faith and believe to be accurate, it is provided for your guidance only. Because many factors may affect processing or application/use, we recommend that you make tests to determine the suitability of a product for your particular prior to use. It does not relieve our customers from obligation to perform a full inspection of the product upon delivery or any other obligation. No warranties of any kind either express or implied, including warranties of merchantability or fitness for a particular purpose are made regarding products described or designs, data or information set forth, or that the products design, data or information may be used without infringing the intellectual property right of others. In no case shall the descriptions, information, data or designs provided be considered a part of our terms and condition of sale.



Ship-to

Analytical Certificate

Delivery 81344664 - 70 **Print date** 2023-07-10 Your reference PO/042604

Our reference Femke den Hartog Material 8571-805 Akoline GC™

Your material no.

Date of shipment 2023-07-07

Batch 4014312181 / Quantity 75 KG / Prod. date 2022-11-29

Inspection lot 3088888

Characteristic	Result		Lower Limit	Target	Upper Limit
Acid value(Syratal LB) Acid value	24.00	mg KOH/g	15.00		35.00
Saponification value(Försåpn Tal LB) Saponification value	247.0	mg KOH/g	225.0		255.0

Recommended shelf life 24 months from production date For detailed information see Product information sheet

> **Quality Control Manager** AAK Sweden AB

This document is electronically produced, and valid without an AAK signature.

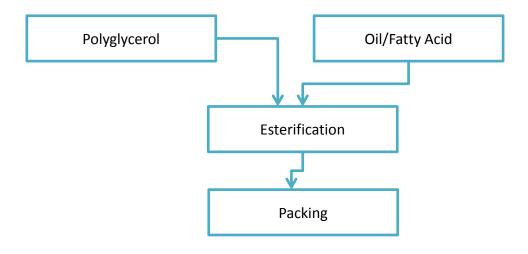
: SE20 5000 0000 0511 8106 1768

Org. no. : 556478-1796 VAT no. : SE556478179601 Approved for Swedish F-tax Registered Office American

A Company in the AAK Group

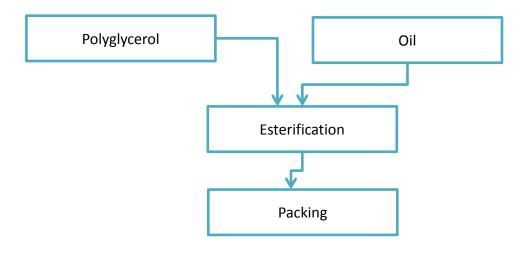
IBAN





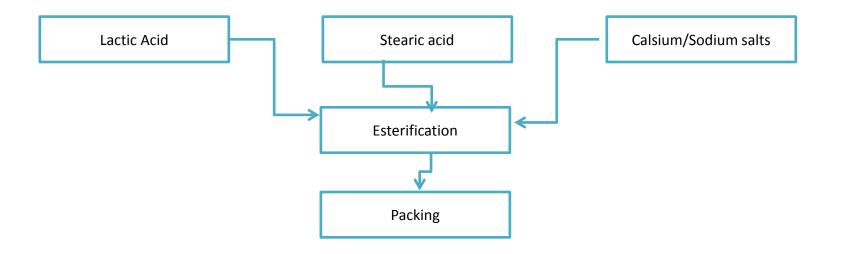
Flowchart Akoline PG7





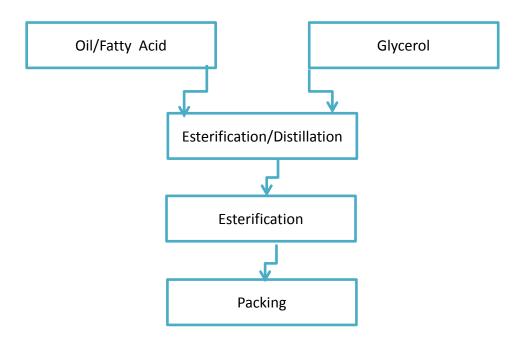
Flowchart Akoline PGPR





Flowchart Akoline SL





Flowchart Akoline LC Akoline GC