Material No 95
Vendor no 86851
Approved by DV
Dorte Vinding

PRODUCT SPECIFICATION – FOOD INGREDIENTS

SALICYLIC ACID PHARM. HUAYIN

BRENNTAG N.V.
BRENNTAG Nederland B.V.

1. PRODUCT INDENTIFICATION

Supplier product name	SALICYLIC ACID PHARM. HUAYIN		
Supplier product number	15617	Date of issue: 25/09/2014	Replaces: 03/07/2013
Origin	Synthesis		
Manufacturer	JQC Huayin		
Country	China		
Certification manufacturer:	ISO 9001		

1.1 Composition

Single ingredient	
Chemical name	Salicylic acid
Chemical formula	$C_7H_6O_3$
Molecular weight	138,12

1.2 Legislative information

CAS-Nº	69-72-7
INTRASTAT CODE	2918210000
EINECS	200-712-3
Legal declaration	

2. PRODUCT INFORMATION

	Unit	Specification	Method
2.1 Physical and Chemical properties			
Appearance		Crystalline powder	
Colour		White to colourless	
Odour/taste		-	
Assay	wt%	99 – 100,5	
Melting point	ºC	158 – 161	
Solubility in water		Slightly soluble	
Solubility in ethanol		Freely soluble	
Solubility in methylene chloride		Sparingly soluble	
2.2 Microbiological data			
N.A.			

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2.3 Contaminants (EC-Regulation 1881/2006)			
2.3.1 Chemical contaminants			
Sulphate (SO_4)	ppm	max. 200	
Loss on drying	%	max. 0,5	
Sulphated ash	%	max. 0,1	
Chloride (Cl)	ppm	max. 100	
Heavy Metals (as Pb)	ppm	max. 20	
Related compounds	%	max. 0,2	
2.3.2 Physical contaminants			
Foreign body control	N.A.		
2.4 Nutritional Information (Directive 90/496/EC, amended by 2003/120/EC and 2008/100/EC)			
2.4.1 Nutritional Values			
Energy	kJ/100g	-	
Energy	kcal/100g	-	
Protein	g/100g	0	
Carbohydrate :	g/100g	0	
Of which Sugars	g/100g	0	
Polyols	g/100g	0	
Starches	g/100g	0	
Others	g/100g	0	
Fat :	g/100g	0	
Of which Saturated	g/100g	0	
Mono-unsaturated	g/100g	0	
Poly-unsaturated	g/100g	0	
Transfatty acids	g/100g	0	
Cholesterol	mg/100g	0	
Water	%	< 1	
Organic acid	g/100g	-	
2.4.2 Minerals			
Sodium	mg/100g		
Others:	mg/100g	-	

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3. FOOD INTOLERANCES

3.1 Allergens (Directives 2000/13/EC, 2003/89/CE 2007/68/EC)

	Y / N	Direct Contamination	Cross-Contamination (Risk)
Barley	N	N	N
Beef	N	N	N
Cacao	N	N	N
Carrot	N	N	N
Celery and celery products	N	N	N
Cereals containing gluten and products produced with these (wheat, rye, oats, spelt, barley)	N	N	N
Chicken	N	N	N
Coriander	N	N	N
Crustaceans and Shellfish	N	N	N
Eggs and egg products	N	N	N
Fish and fish products	N	N	N
Glutamate	N	N	N
Lupin and products thereof	N	N	N
Milk and milk products (incl. Lactose)	N	N	N
Molluscs and products thereof	N	N	N
Mustard and mustard products	N	N	N
Nuts and nut products (almonds, hazelnuts, walnuts)	N	N	N
Peanuts and peanut products	N	N	N
Pork	N	N	N
Sesame and sesame products	N	N	N
Soybean and soybean products	N	N	N
Sulphite (E221 - E228)	N	N	N
Sulphur dioxide (> 10 mg/kg)	N	N	N

3.2 Suitability for other diets

Coeliacs	Y	Lactose intolerant	Y
Halal	N	Vegans	Y
Kosher	N	Vegetarian	Y

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3.3 GMO Declaration, acc. Regulations 298/2008/EC and 1830/2003/EC

SALICYLIC ACID PHARM. does not contain genetically modified organisms and is not produced using raw materials of a genetically modified origin. At no stage during production does the product comes into contact with genetically modified organisms.
SALICYLIC ACID PHARM. is falling under the scope of the EC Regulation n° 1829/2003 on genetically modified food and feed and the EC Regulation n° 1830/2003 on the traceability and labeling of genetically modified organisms and the traceability of products derived from them and to modification of Directive 2001/18/EC.

3.4 Irradiation, acc. Directive 1999/2/EC and 1999/3/EC

This product was not subject to any kind of irradiation treatment

3.5 BSE/TSE Declaration

The used ingredients for SALICYLIC ACID PHARM. are not of animal origin. The processing equipment and the packing material which is used to manufacture, pack or fill the products into the packing units do not come into contact with any meat or meat-by product.

3.6 Residual Solvents, complies with

For the manufacturing of the product, no solvents are used. This product is in compliance with the Guideline CPMP/ICH/283/95 and chapter <467> of the current edition of the USP-NF for residual solvents.

3.7 Regulatory information

The product complies with:

BP-98

4. PACKAGING

Packaging description	Bags
Packaging net content	25 kg
Different packagings on request	Yes
Paletisation	1000 kg

5. STORAGE CONDITIONS

Storage conditions	Must be kept in a cool, dry and well ventilated environment, in closed original packaging
Shelf life (recommended re-analysis)	36 months after production, under the above mentioned conditions

6. DISCLAIMER

The content of the Product Specification Sheet is completed to the best of our knowledge.
This document does not dismiss the user of his legal obligation with respect to pharma safety.

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COMPANY INFORMATION DISTRIBUTOR		
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country	Belgium	The Netherlands
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website	www.brenntag.be	www.brenntag.nl
e-mail	info@brenntag.be	info@brenntag.nl
activities	Distribution and export of chemicals and raw materials	
VAT number	BE0405317567	NL001375945B01
recall procedure available	Yes	
emergency number (24/365)	+32 (0)56 77 69 44	+31 (0)78 6544 944
QUALITY SYSTEMS		
ISO 9001	Yes	Yes
ISO 14001	Yes	Yes
ISO 22000	Yes	Yes
FSSC 22000	Yes	Yes
GMP+ -feed	Yes	Yes
OHSAS18001	-	Yes
ESAD	Yes	Yes
other	-	AEO