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DATA SHEET

EUCALYPTUS G 80/85 ORG OIL GMP

Eucalyptus globulus Labillardiere

Number of pages : 2

 Created on
 : 02/11/2003

 Modified on
 : 18/06/2019

 Last update
 : 08/03/2022

TDS B381P

Version 04.55

Organic product certified by FR-BIO-01

PROCESSING METHOD

Essential Oil obtained by steam distillation from fresh leaves or fresh terminal stems of Eucalyptus globulus Labillardiere

PHYSICAL PROPERTIES

Odour : Camphorated, fresh and note of cineol

Appearance : Liquid

Color : Colorless to pale yellow

CHEMICAL PROPERTIES

Specific gravity at 20° C : 0,906 to 0,927 Refractive index at 20° C : 1,458 to 1,470 Optical rotation at 20° C : +0,0 ° to +10,0 °

Flash point : +45 ℃
Acid value : N.A.

Peroxide value : N.A.

MAIN INGREDIENTS

Cineol 1,8 (Eucalyptol) (70,00 to 87,00%)

D-Limonene (0,05 to 15,00%) Alpha pinene (0,05 to 10,00%)

Alpha phellandrene (0,05 to 1,50%)

Beta pinene (0,05 to 1,50%)

Sabinene (<= 0,30%)

Camphor (<= 0,10%)

TECHNICAL SPECIFICATIONS

NCS, Natural Complex Substance (100% pure and natural)

N°CAS TSCA : **8000-48-4** ; **1377456-89-7**

N°CAS EINECS : 84625-32-1 **N°EINECS** : 283-406-2 N°FEMA : 2466 N°FDA : 172.510 N°CoE: : 185n FCC : X **RIFM** : X **FMA** : X

AFNOR : **NF T 75-225**

INCI : EUCALYPTUS GLOBULUS LEAF OIL

Country of harvest : PORTUGAL

Country of : PORTUGAL

manufacturing

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HANDLING, STORING AND SHELF LIFE

Follow usual precaution of chemical ingredient handling. Store in a sealed container, in a dry, cool and aired area, protected from the light.

Shelf life: 24 months from the date of manufacture

TRANSPORT INFORMATION

Hazard Class 3, Packing group III, UN n°1169

Customs rate code : 3301 29 49

SPECIAL INDICATIONS

■ The presence of the following allergen substances in a finished product must be indicated by way of labelling if their respective concentration exceeds 100 ppm in a rinsed product and 10 ppm in a product not rinsed. (Annex III of Regulation (EC) N°1223/2009 of 30 november 2009 on cosmetic products)

Present allergens : D-Limonene (0,05 to 15,00%), Geraniol (<= 0,50%)

■ IFRA restrictions : This substance and/or some of its components are covered by the Code of Practice of the IFRA in

effect, available on the internet website www.ifraorg.org

Comment: Regarding the number of significant figures of the specifications, it is useful to refer to both, COA of the batch and pharmacopoeia rules.

Contact M. DENAT Jean-François

This sheet has been updated on 08/03/2022

Number of pages: 2 Printed on: 08/03/2022 End of document

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