MARK. HELP WANTED ANTI-ACNE EXFOLIATING CLEANSER- salicylic acid paste Avon Products, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

Active ingredient

Salicylic Acid 0.5%.....

Purpose

.....Acne Treatment

Uses

- for the treatment of acne
- helps clear up acne blemishes
- helps prevent new acne blemishes from forming
- penetrates pores to eliminate most blackheads and acne pimples

Warnings

For external use only

When using this product

- avoid contact with eyes. If product contacts eyes, rinse thoroughly with water. If irritation persists, seek medical attention.
- skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical medication at a time.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- wet hands and face with warm water
- pour into hands and work into a lather
- gently massage into skin in a circular motion, avoiding the eye area
- rinse and pat dry.
- because excessive drying of the skin may occur, start with one application daily, then gradually increase to two to three times daily if needed or as directed by a doctor
- if bothersome dryness, irritation or peeling occurs, reduce application to every other day

Inactive ingredients: WATER/EAU

SODIUM LAURETH SULFATE

KAOLIN

COCAMIDE MEA

GLYCOL STEARATE

SODIUM COCO-SULFATE

HYDROGENATED CASTOR OIL

PYRUS MALUS (APPLE) JUICE

VANILLA PLANIFOLIA FRUIT EXTRACT

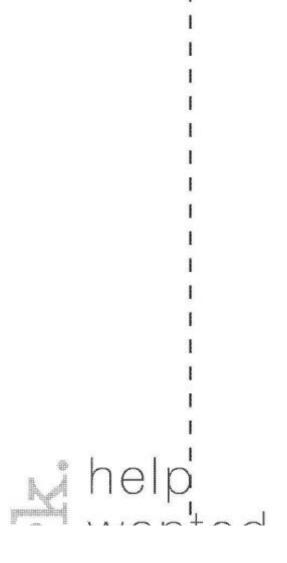
ROSMARINUS OFFICINALIS (ROSEMARY) LEAF EXTRACT

MENTHA PIPERITA (PEPPERMINT) LEAF EXTRACT

CINNAMOMUM CASSIA BARK EXTRACT CUCUMIS SATIVUS (CUCUMBER) FRUIT EXTRACT PEG-40 HYDROGENATED CASTOR OIL **POLYETHYLENE BUTYLENE GLYCOL** SHEA BUTTERAMIDOPROPYL BETAINE SD ALCOHOL 40-B **METHYLPARABEN** DISODIUM EDTA GLYCINE SOJA (SOYBEAN) OIL HELIANTHUS ANNUUS (SUNFLOWER) SEED OIL HYDRATED SILICA PHOSPHORIC ACID CAPRYLIC/CAPRIC TRIGLYCERIDE ISOPROPYL MYRISTATE DIPROPYLENE GLYCOL HEXYLENE GLYCOL **METHYLCHLOROISOTHIAZOLINONE METHYLISOTHIAZOLINONE** TITANIUM DIOXIDE/CI 77891

Questions?

Call 1-800-MEET-MARK





150 ml 5 FL OZ

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Inactive ingredients: Water/Eau, Sodium Laureth Sulfate, Kaolin, Cocamide Mea, Glycol Stearate, Sodium Coco-Sulfate, Hydrogenated Castor Oil, Pyrus Malus (Apple) Juice, Vanilla Planifolia Fruit Extract, Rosmarinus Officinalis (Roserhary) Leaf Extract, Mentha Piperita (Peppermint) Leaf Extract, Cirinamomum Cassia Bark Extract, Cucumis Sativus (Cucumber) Fruit Extract, PEG-40 Hydrogenated Castor Oil, Polyethylene, Butylene Glycol, Shea Butteramidopropyl Betaine, SD Alcohol 40-B, Methylparaben, Disodium EDTA, Glycine Soja (Soybean) Oil, Helianthus Annuus (Sunflower) Seed Oil, Hydrated Silica, Phosphoric Acid, Caprylic/Capric Triglyceride, Isopropyl Myristate, Dipropylene Glycol, Hexylene Glycol, Methylchloroisothiazolinone, Methylisothiazolinone, Titanium Dioxide/Cl 77891.

Questions? Call 1-800-MEET-MARK

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salicylic acid paste

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10096-0237
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.75 mL in 150 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
SO DIUM LAURETH SULFATE (UNII: BPV390 UAP0)		
KAOLIN (UNII: 24H4NWX5CO)		

GLYCOL STEARATE (UNII: 0324G66D0E)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
METHYLPARABEN (UNII: A2I8 C7HI9 T)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
PHO SPHORIC ACID (UNII: E4GA8884NN)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
HEXYLENE GLYCOL (UNII: KEH0 A3F75J)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229 D0 E1QFA)	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10096-0237-1	150 mL in 1 TUBE		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	12/20/20 10	

Labeler - Avon Products, Inc. (001468693)

Establishment			
Name	Address	ID/FEI	Business Operations
Avon Products, Inc.		005149471	manufacture

Revised: 12/2010 Avon Products, Inc.