PHISODERM CLEAN ANTI-BLEMISH FACIAL CLEANSER- salicylic acid gel The Mentholatum Company

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 2%

Purpose

Acne treatment

Uses

For the treatment of acne

Warnings

For external use only

When using this product

- avoid contact with eyes. If contact occurs, flush thoroughly with water.
- 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 acne medication at a time.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

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

Directions

- wet face, then gently massage cleanser onto skin
- rinse well and pat dry
- use twice daily

Inactive ingredients

water, lauramidopropyl betaine, sodium C14-16 olefin sulfonate, lauryl glucoside, aloe

barbadensis leaf juice, aminomethyl propanol, butylene glycol, caprylyl/capryl glucoside, carica papaya (papaya) fruit extract, chamomilla recutita (matricaria) flower extract, disodium EDTA, fragrance, glycerin, hydroxyphenyl propamidobenzoic acid, hydroxypropyltrimonium hyaluronate, pentylene glycol, potassium sorbate, sodium benzoate, sodium chloride, tocopheryl acetate [vitamin E]

Questions?

1-877-636-2677 MON-FRI 9 AM to 5 PM (EST)

Principal Display Panel



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PHISODERM CLEAN ANTI-BLEMISH FACIAL CLEANSER

salicylic acid gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10742-1235	

Route of Administration TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20 mg in 1 mL		

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
LAURAMIDOPROPYL BETAINE (UNII: 23D6XVI233)			

SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)	
PAPAYA (UNII: KU94FIY6JB)	
CHAMOMILE (UNII: FGL3685T2X)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYPHENYL PROPAMIDOBENZOIC ACID (UNII: 25KRT26H77)	
DIHYDROXYPROPYLTRIMONIUM (UNII: 925G44QFFO)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:10742- 1235-1	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/02/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	07/02/2021	

Labeler - The Mentholatum Company (002105757)

Registrant - The Mentholatum Company (002105757)

Establishment					
Name	Address	ID/FEI	Business Operations		
The Mentholatum Company		002105757	manufacture(10742-1235)		

Revised: 2/2023 The Mentholatum Company