PHISODERM ANTI-BLEMISH GEL 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

treats and helps prevent acne blemishes

Warnings

For external use only

When using this product

- avoid contact with the 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 onto skin
- Rinse well and pat dry
- use twice daily

Inactive ingredients

water, lauramidopropyl betaine, sodium C14-16 olefin sulfonate, lauryl glucoside, trolamine, butylene glycol, chamomilla recutita (matricaria) flower extract, disodium EDTA, DMDM hydantoin, fragrance, glycerin, hydroxyphenyl propamidobenzoic acid, iodopropynyl butylcarbamate, lactic acid, pentylene glycol, polysorbate 20, sodium chloride

Questions?

Toll free **1-877-636-2677** MON-FRI 9 AM-5 PM (EST)

Principal Display Panel



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The Mentholatum Company, Orchard Park, NY 14127 @2013 www.mentholatum.com





PHISODERM ANTI-BLEMISH GEL CLEANSER

salicylic acid gel

Product Information

Route of Administration

Product Type HUMAN OTC DRUG **Item Code (Source)** NDC:10742-8417 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 g	

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
LAURAMIDOPROPYL BETAINE (UNII: 23D6XVI233)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
TROLAMINE (UNII: 903K93S3TK)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CHAMOMILE (UNII: FGL3685T2X)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYPHENYL PROPAMIDOBENZOIC ACID (UNII: 25KRT26H77)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
LACTIC ACID, UNSPECIFIED FORM (UNII: 33X04XA5AT)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742- 8417-1	177 g in 1 TUBE; Type 0: Not a Combination Product	09/01/2013	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part333D	09/01/2013		

Labeler - The Mentholatum Company (002105757)

Registrant - The Mentholatum Company (002105757)

Establishment				
Name	Address	ID/FEI	Business Operations	
The Mentholatum Company		002105757	manufacture(10742-8417)	

Revised: 2/2023 The Mentholatum Company