PHISODERM CLEAR CONFIDENCE DAILY GEL FACE WASH- salicylic acid liquid
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, sodium C14-16 olefin sulfonate, lauramidopropyl betaine, lauryl glucoside, sodium chloride, disodium EDTA, DMDM hydantoin, fragrance, iodopropynyl butylcarbamate, lactic acid, polysorbate 20, triethanolamine

Questions?

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

Principal Display Panel


pHisoderm®
clear
confidence®
Daily
Gel Face Wash
clears up blemishes for a clean, healthy complexion
2% salicylic acid
acne treatment
Oil-Free
gently cleans without over drying
6 FL OZ (177 mL)
LB593001

Principal Display Panel

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PHISODERM CLEAR CONFIDENCE DAILY GEL FACE WASH

salicylic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10742-1412
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)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
LAURAMIDOPROPYL BETAINE (UNII: 23D6XVI233)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
LACTIC ACID, UNSPECIFIED FORM (UNII: 33X04XA5AT)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:10742-1412-1	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	05/01/2012	

Labeler - The Mentholatum Company (002105757)

Registrant - The Mentholatum Company (002105757)

Establishment

Name	Address	ID/FEI	Business Operations
The Mentholatum Company		002105757	manufacture(10742-1412)

Revised: 2/2023

The Mentholatum Company