

**PHISODERM CLEAR CONFIDENCE DAILY GEL FACE WASH- salicylic acid liquid**  
**The Mentholatum Company**

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**Drug Facts - pHisoderm Clear Confidence Daily Gel Face Wash**

**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





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The Mentholatum Company, Orchard Park, NY 14127 © 2012



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# PHISODERM CLEAR CONFIDENCE DAILY GEL FACE WASH

salicylic acid liquid

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:10742-1412
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>SALICYLIC ACID</b> (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM C14-16 OLEFIN SULFONATE</b> (UNII: O9W3D3YF5U)	
<b>LAURAMIDOPROPYL BETAINE</b> (UNII: 23D6XVI233)	
<b>LAURYL GLUCOSIDE</b> (UNII: 76LN7P7UCU)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>DMDM HYDANTOIN</b> (UNII: BYR0546TOW)	
<b>IODOPROPYNYL BUTYLCARBAMATE</b> (UNII: 603P14DHEB)	
<b>LACTIC ACID, UNSPECIFIED FORM</b> (UNII: 33X04XA5AT)	
<b>POLYSORBATE 20</b> (UNII: 7T1F30V5YH)	
<b>TROLAMINE</b> (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 Drug	M006	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: 12/2024

The Mentholatum Company