

**OXY VOLCANIC ASH ACNE 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.*

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**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**

- keep away from 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 gently
- gently massage cleanser onto skin, avoiding eye area
- rinse thoroughly

**Inactive ingredients**

acrylates copolymer, aloe barbadensis leaf juice, anhydrous citric acid, butylene glycol,

charcoal powder, cocamidopropyl betaine, edetate disodium, fragrance, glycerin, menthol, portulaca oleracea extract, potassium C12-13 alkyl phosphate, potassium sorbate, purified water, silica, sodium benzoate, sodium C14-16 olefin sulfonate, trolamine, volcanic ash

## Questions?

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

## Package/Label Principal Display Panel



## Principal Display Panel



## OXY VOLCANIC ASH ACNE FACE WASH

salicylic acid liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:10742-8334
<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>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	

<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)
<b>ACTIVATED CHARCOAL</b> (UNII: 2P3VMU3H10)
<b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)
<b>MENTHOL, UNSPECIFIED FORM</b> (UNII: L7T10EIP3A)
<b>PURSLANE</b> (UNII: M6S840WVG5)
<b>POTASSIUM SORBATE</b> (UNII: 1VPU26JZZ4)
<b>WATER</b> (UNII: 059QF0KO0R)
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)
<b>SODIUM C14-16 OLEFIN SULFONATE</b> (UNII: O9W3D3YF5U)
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-8334-1	207 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2018	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	11/01/2018	

**Labeler** - The Mentholatum Company (002105757)

**Registrant** - The Mentholatum Company (002105757)

### Establishment

Name	Address	ID/FEI	Business Operations
The Mentholatum Company		002105757	manufacture(10742-8334)

Revised: 2/2023

The Mentholatum Company