

OXY VOLCANIC ASH ACNE FACE WASH- salicylic acid liquid
The Mentholatum Company

Drug Facts - OXY Volcanic Ash Acne 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

- 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

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

BUTYLENE GLYCOL (UNII: 3XUS85K0RA)
ACTIVATED CHARCOAL (UNII: 2P3VMU3H10)
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)
EDETATE DISODIUM (UNII: 7FLD91C86K)
GLYCERIN (UNII: PDC6A3C0OX)
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)
PURSLANE (UNII: M6S840WYG5)
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)
WATER (UNII: 059QF0K00R)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
SODIUM BENZOATE (UNII: OJ245FE5EU)
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)
TROLAMINE (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 Drug	M006	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: 12/2024

The Mentholatum Company