OXY VOLCANIC ASH DETOX SCRUB- salicylic acid cream
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

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

bentonite, butylene glycol, ceteareth-20, cetearyl alcohol, cetostearyl alcohol, charcoal

powder, cocamidopropyl betaine, cocos nucifera (coconut) oil, disodium cocoamphodiacetate, edetate disodium, FD&C blue no. 1 aluminum lake, FD&C red no. 40 aluminum lake, FD&C yellow no. 6 aluminum lake, fragrance, glycerin, hydroxypropyl starch phosphate, jojoba esters, kaolin, lecithin, menthol, methyl gluceth-20, panthenol, paraffin, potassium sorbate, purified water, silica, sodium benzoate, sodium laureth sulfate, trolamine, volcanic ash, xanthan gum

Questions?

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

Package/Label Principal Display Panel

OXY VOLCANIC ASH DETOX SCRUB
salicylic acid cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10742-1338
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 g

Inactive Ingredients

Ingredient Name	Strength
BENTONITE (UNII: A3N5ZCN45C)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWJY)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
ACTIVATED CHARCOAL (UNII: 2P3VWU3H10)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
COCONUT OIL (UNII: Q9L0O73W7L)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGENATED JOJOBA OIL, RANDOMIZED (UNII: Q47ST02F58)	
KAOLIN (UNII: 24H4NWX5CO)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
PANTHENOL (UNII: WV9CM0O67Z)	
PARAFFIN (UNII: I9O0E3H2ZE)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
WATER (UNII: 059QF0KO0R)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
TROLAMINE (UNII: 9O3K93S3TK)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-1338-1	142 g in 1 TUBE; 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(10742-1338)

Revised: 2/2023 The Mentholatum Company