

OXY TOTAL CARE CLARIFYING MOISTURIZER- benzoyl peroxide lotion
The Mentholatum Company

Drug Facts - OXY Total Care Clarifying Moisturizer

Active ingredient

Benzoyl peroxide 2.5%

Purpose

Benzoyl peroxide - Acne treatment

Keep Out of Reach of Children

If swallowed, get medical help or contact a Poison Control Center right away.

Uses

- For the treatment of acne.
- Helps prevent new acne blemishes.

Warnings

For external use only.

Do not use if you

- have very sensitive skin
- are sensitive to benzoyl peroxide.

When using this product

- skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, use only one topical acne medication at a time.
- avoid unnecessary sun exposure and use a sunscreen
- avoid contact with the eyes, lips, and mouth
- avoid contact with hair and dyed fabrics, which may be bleached by this product
- skin irritation may occur, characterized by redness, burning, itching, peeling, or possibly swelling. Irritation may be reduced by using the product less frequently or in a lower concentration.

Stop use and ask a doctor if irritation becomes severe.

If pregnant or breast-feeding, ask a health professional before use.

Directions

- clean the skin thoroughly before applying this product

- use a dime-sized amount to cover the entire affected area with a thin layer one to two times daily
- because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor
- if bothersome dryness or peeling occurs, reduce application to once a day or every other day.
- if going outside, apply sunscreen after using this product. If irritation or sensitivity develops, stop use of both products and ask a doctor.
- *Sensitivity Test for a New User.* Apply product sparingly to one or two small affected areas during the first 3 days. If no discomfort occurs, follow the directions stated above.

Other information

- THIS PRODUCT MAY BLEACH HAIR OR DYED FABRICS
- KEEP TIGHTLY CLOSED
- avoid storing at temperatures above 100°F (38°C)

Inactive Ingredients

water, glycerin, butylene glycol, carbomer, chlorphenesin, citric acid, dimethyl isosorbide, disodium EDTA, hyaluronic acid, hydrolyzed algin, lauryl methacrylate/glycol dimethacrylate crosspolymer, PEG/PPG-4/12 dimethicone, phenoxyethanol, phytic acid, portulaca oleracea extract, silanetriol, sodium citrate, sodium hydroxide, xanthan gum, zinc sulfate

Questions?1-877-636-2677

MON-FRI 9AM-5PM (EST)

Package/Label Principal Display Panel



OXY TOTAL CARE CLARIFYING MOISTURIZER

benzoyl peroxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10742-1314
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOYL PEROXIDE (UNII: W9WZ N9A0GM) (BENZOYL PEROXIDE - UNII:W9WZ N9A0GM)	BENZOYL PEROXIDE	25 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
EDETATE DISODIUM (UNII: 7FLD91C86K)	
HYALURONIC ACID (UNII: S270N0TRQY)	
ALGINIC ACID (UNII: 8C3Z4148WZ)	
LAURYL METHACRYLATE/GLYCOL DIMETHACRYLATE CROSSPOLYMER (UNII: EX0F4CZ66H)	
PEG/PPG-4/12 DIMETHICONE (UNII: JAN3585W85)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
FYTIC ACID (UNII: 7IGF0S7R8I)	
PURSLANE (UNII: M6S840WYG5)	
SILANETRIOL (UNII: E52D0J3TS5)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ZINC SULFATE HEPTAHYDRATE (UNII: N57JI2K7WP)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-1314-1	1 in 1 CARTON	09/30/2019	
1		100 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	09/30/2019	

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

Establishment

Name	Address	ID/FEI	Business Operations
The Mentholatum Company		002105757	manufacture(10742-1314)

Revised: 12/2024

The Mentholatum Company