

OXY SENSITIVE FACE WASH MAXIMUM SOOTHING- benzoyl peroxide lotion

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

Benzoyl peroxide 5%

Purpose

Acne treatment

Uses

treats and helps prevent acne

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, only use 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 this 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.

Keep Out of Reach of Children

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

Directions

- wet face
- apply to hands then work into a lather and massage gently onto face
- rinse thoroughly and pat dry
- because excessive drying of the skin may occur, start with 1 use daily, then gradually increase to 2 or 3 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:* Apply product sparingly to a small affected area for the first 3 days. If no discomfort occurs, follow directions 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, cetostearyl alcohol, sodium C14-16 olefin sulfonate, disodium laureth sulfosuccinate, capryl/capramidopropyl betaine, xanthan gum, butylene glycol, citric acid, diazolidinyl urea, hydrolyzed soy protein, methylparaben, PEG-8 dimethicone, portulaca oleracea extract, propylparaben, sodium citrate, sodium hydroxide, sodium lauroyl sarcosinate

Questions?

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

Principal Display Panel

NEW LOOK!
Coming Soon

ACNE CLEANSER
Sensitive Skin

5%
Benzoyl Peroxide
Acne Treatment

OXY
ACNE MEDICATION

ACNE CLEANSER
Sensitive Skin

5% Benzoyl Peroxide Acne Treatment

Eliminates
Acne Bacteria
Fragrance Free

5 FL OZ (148 mL)

Drug Facts Clinically Proven Active Ingredient

Active ingredient	Purpose
Benzoyl peroxide 5%	Acne treatment

Uses treats and helps prevent acne

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AC017003

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OXY SENSITIVE FACE WASH MAXIMUM SOOTHING

benzoyl peroxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10742-1313
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	50 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
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WATER (UNII: 059QF0KO0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
DISODIUM LAURETH SULFO SUCCINATE (UNII: D6DH1DTN7E)	
CAPRYL/CAPRAMIDOPROPYL BETAINE (UNII: 231H3ZT9NE)	
XANTHAN GUM (UNII: TTV12P4NEE)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
SOY PROTEIN (UNII: R44IWB3RN5)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
PEG-8 DIMETHICONE (UNII: GIA7T764OD)	
PURSLANE (UNII: M6S840WXG5)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM LAUROYL SARCOSINATE (UNII: 632GS99618)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-1313-1	148 mL in 1 TUBE; Type 0: Not a Combination Product	08/01/2016	
2	NDC:10742-1313-2	185 mL in 1 TUBE; Type 0: Not a Combination Product	08/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	08/01/2016	

Labeler - The Mentholatum Company (002105757)

Revised: 11/2019

The Mentholatum Company