OXY ADVANCED CARE RAPID SPOT TREATMENT- benzoyl peroxide gel
The Mentholatum Company

Drug Facts

Active ingredient

Benzoyl peroxide 10%

Purpose

Acne treatment

Uses

- for the treatment of acne
- helps prevent new acne blemishes from forming

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 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.

Keep out of reach of children.

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

Directions

- clean the skin thoroughly before applying this product
- cover the entire affected area with a thin layer 1 to 3 times daily
- because excessive drying of the skin may occur, start with 1 application 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.

Other Safety Information

- avoid storing at temperatures above 100 ^{O}F (38 ^{O}C)

Inactive Ingredients

water, glycerin, butylene glycol, carbomer, chlorphenesin, citric acid, dimethyl isosorbide, disodium EDTA, fragrance, hydrated silica, hydrolyzed soy protein, lauryl methacrylate/glycol dimethacrylate crosspolymer, PEG/PPG-4/12 dimethicone, phenoxyethanol, portulaca oleracea extract, propanediol, rhodomyrtus tomentosa fruit extract, sodium citrate, sodium hydroxide, xanthan gum

Questions

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

Package/Label Principal Display Panel

MENTHOLATUM

2.250 X 1.125 X 5.000 - PRINTSIDE

S200263 - 3/16/20

ART #:_____

ARTIST:_____

PROOFER:_____

PLATE #:_____

PLATE LOC:_____

☐ KO's REQUIRED ☐ NO COATING ☐ GLOSS UV COATING
☐ HOT STAMP ☐ UV FLOOD ☐ MATTE UV STRIKE THRU VARNISH
☐ COLD FOIL ☐ EMBOSS ☐ _____

BARCODE VERIFICATION:
☐ UPC ☐ EAN/128
☐ 2 of 5 ☐ QR Code
☐ Other: _____

COATING KO

THIS PROOF WAS PRODUCED WITH G7® TECHNOLOGY

C CYAN M MAGENTA Y YELLOW B BLACK X PMS 360 Z U V DIELINE

OXY ADVANCED CARE RAPID SPOT TREATMENT

benzoyl peroxide gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10742-1201
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	100 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
SOY PROTEIN (UNII: R44IWB3RN5)	
LAURYL METHACRYLATE/GLYCOL DIMETHACRYLATE CROSSPOLYMER (UNII: EX0F4CZ66H)	
PEG/PPG-4/12 DIMETHICONE (UNII: JAN3585W85)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PURSLANE (UNII: M6S840WXG5)	
PROPANEDIOL (UNII: 5965N8W85T)	
RHODOMYRTUS TOMENTOSA FRUIT (UNII: Q99511S58K)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:10742-1201-1	1 in 1 CARTON	07/12/2021	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:10742-1201-2	1 in 1 CARTON	12/01/2021	
2		32.6 g in 1 TUBE; Type 0: Not a Combination Product		

3	NDC:10742-1201-3	1 in 1 CARTON	02/21/2022	
3		18.4 g 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	07/12/2021	

Labeler - The Mentholatum Company (002105757)

Revised: 2/2024 The Mentholatum Company