BENZOYL PEROXIDE- benzoyl peroxide gel Malezia LLC

Benzoyl Peroxide

50 mL NDC: 77924-002-01



BIOCHEMICALLY CLEARER SKIN

50 ML I 1.7 FL OZ

Drug Facts

Active ingredient

Purpose

Benzoyl Peroxide 2.5% Acne Treatment

Use For the treatment of acne.

Warnings:

•For external use only.

Do not use if you

- Have very sénsitive skin.
- Are sensitive to benzoyl peroxide.
 When using this product
- Avoid unnecessary sun exposure and use an SPF.
- 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.

- Only use one topical acne medication at a time as skin irritation and dryness is more likely to occur.
- Keep out of reach of children.
- If swallowed get medical help or call poison control center immediately.

Stop use and ask a doctor if

Irritation becomes severe.

Directions:

- Clean the skin gently before applying this product.
- Cover the affected area with a fingertip unit of product once or twice daily.
- If bothersome dryness or peeling occurs, reduce application to once a day or every other day.

Other Ingredients: Water, Propanediol, Glycerin, Carbomer, Sodium Phytate, Sodium Hydroxide

Malezia.com
Spokane Valley, WA 99216
Designed in the USA
Made in PRC





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BENZOYL PEROXIDE benzoyl peroxide gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:77924-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of
Strength

Strength

BENZOYL PEROXIDE 25 mg in 1000 mg

Inactive Ingredients			
Ingredient Name	Strength		
GLYCERIN (UNII: PDC6A3C0OX)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			
WATER (UNII: 059QF0KO0R)			
CARBOMER 980 (UNII: 4Q93RCW27E)			
PHYTATE SODIUM (UNII: 88496G1ERL)			

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

ı	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:77924-002- 02	50000 mg in 1 TUBE; Type 0: Not a Combination Product	12/31/2020	

Marketing Information				
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M006	08/21/2020		

Labeler - Malezia LLC (123312045)

PROPANEDIOL (UNII: 5965N8W85T)

Establishment				
Name	Address	ID/FEI	Business Operations	
Malezia LLC		123312045	label(77924-002)	

Malezia LLC Revised: 4/2024