

24 HOUR ACNE SERUM - benzoyl peroxide gel
Drmtlgy, LLC

DRMTLGY® 24 Hour Acne Serum

Drug Facts

Active ingredient

Benzoyl Peroxide 5%

Purpose

Acne Medication

Use

For the treatment of acne.

Warnings

For external use only.

When using this product • avoid unnecessary sun exposure and use sunscreen. • avoid contact with eyes, lips and mouth. • avoid contact with hair and dyed products, 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

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. Apply affected area with a thin layer, avoiding eye area. Allow to absorb before applying additional products. Can be used twice daily or as directed by a physician. If irritation or sensitivity develops, stop use of product and consult a physician.

Inactive Ingredients

Water, Glycolic Acid, Sclerotium Gum, Arnica Montana Flower Extract, Allantoin, Echinacea Purpurea Extract, Hydrastis Canadensis (Goldenseal) Extract, Lavandula Angustifolia (Lavender) Flower/Leaf/Stem Extract, Calendula Officinalis Flower Extract, Glycerin, Gluconolactone, Sodium Benzoate, Tetrasodium EDTA, Sodium Hydroxide

Questions or comments?

1-888-DRMTLGY

MEDICAL GRADE SKIN CARE

REPAIR

clear//prevent

DRMTLGY, LLC
Chatsworth, CA 91311
www.DRMTLGY.com

Made in USA

Packaging



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ACNE SERUM

REPAIR
clear//prevent

1.1 oz / 32 g

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24 HOUR ACNE SERUM

benzoyl peroxide gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83286-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)		BENZOYL PEROXIDE	5 g in 100 g	
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
GLYCOLIC ACID (UNII: 0WT12SX38S)				
BETASIZOFIRAN (UNII: 2X51AD1X3T)				
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)				
ALLANTOIN (UNII: 344S277G0Z)				
ECHINACEA PURPUREA WHOLE (UNII: QI7G114Y98)				
GOLDENSEAL (UNII: ZW3Z11D0JV)				
LAVANDULA ANGUSTIFOLIA SUBSP. ANGUSTIFOLIA FLOWERING TOP (UNII: 9YT4B71U8P)				
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)				
GLYCERIN (UNII: PDC6A3C0OX)				
GLUCONOLACTONE (UNII: WQ29KQ9POT)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
EDETATE SODIUM (UNII: MP1J8420LU)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
Product Characteristics				
Color	white (White to Off White)	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83286-007-01	32 g in 1 BOTTLE; Type 0: Not a Combination Product	04/08/2025	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M006	04/08/2025	

Labeler - Drmtlgy, LLC (094762235)