ACNE CONTROL SERUM- benzoyl peroxide lotion
Private Label Skin Care

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.

Acne Control 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 physician. If irritation or sensitivity develops, stop use of product and consult physician.

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

brand MD®

SKIN CARE

ACNE DEFENSE

medical grade

✓ Clinically proven to decrease sebum production by up to 70%

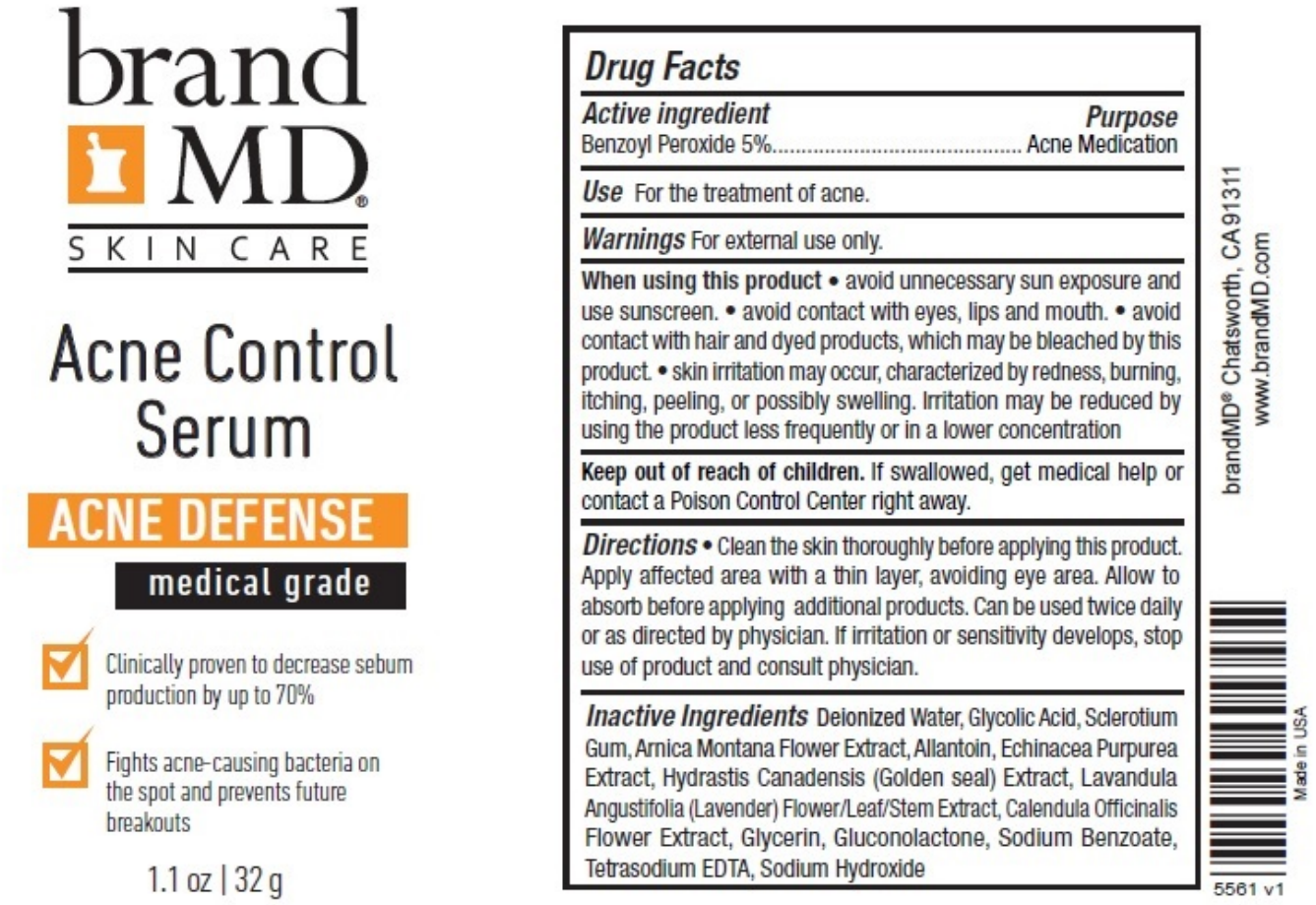
✓ Fights acne-causing bacteria on the spot and prevents future breakouts

brandMD® Chatsworth, CA 91311

www.brandMD.com

Made in USA

Packaging



ACNE CONTROL SERUM

benzoyl peroxide lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72957-001(NDC:39765-030)
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72957-001-01	32 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	01/23/2020	

Labeler - Private Label Skin Care (116996962)

Establishment

Name	Address	ID/FEI	Business Operations
Private Label Skin Care		116996962	relabel(72957-001)

Revised: 1/2022 Private Label Skin Care