

NEUTROGENA STUBBORN ACNE AM TREATMENT- benzoyl peroxide gel
Kenvue Brands LLC

Neutrogena® STUBBORN ACNE AM TREATMENT

Drug Facts

Active ingredient

Benzoyl Peroxide (2.5%)

Purpose

Acne treatment

Use

For the treatment of acne.

Warnings

For external use only.

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. If contact occurs, flush thoroughly with water
- avoid contact with hair or 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.

Do not use if you

- have very sensitive skin
- are sensitive to benzoyl peroxide.

Stop use and ask a doctor if

- Irritation becomes severe.

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 one to three times daily
- because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three 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 Information

- Store at 20°C to 25°C (68°F - 77°F)

Inactive ingredients

Water, Carbomer Homopolymer type B, Ethylhexylglycerin, Sodium Hydroxide, Chlorphenesin, Disodium EDTA, Laureth-4, Hydroxypropyl Methylcellulose

Questions?

800-582-4048; Outside US, dial collect **215-273-8755** or visit www.neutrogena.com

Distributed by:

**JOHNSON & JOHNSON
CONSUMER INC.**

Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 56 g Tube Carton

New

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

STUBBORN

ACNE

AM TREATMENT

Benzoyl Peroxide

Acne treatment

**works all day to help
eliminate stubborn acne**

2.5% Micronized BPO

NET WT. 2.0 OZ

(56 g)

Neutrogena
**STUBBORN
ACNE™**
AM TREATMENT ☀️

NEW

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DERMATOLOGIST RECOMMENDED BRAND

**STUBBORN
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Acne Treatment

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(56 g)



Our recommended regimen
for stubborn acne
and post-acne marks:



**Stubborn Acne
AM Treatment**



**Stubborn Marks
PM Treatment**

When used together, the Retinol
helps release pore-clogging dead
skin cells, giving Benzoyl Peroxide a
clear path to effectively target acne-
causing bacteria.



**2.5% Micronized
Benzoyl Peroxide**

A recommended first line of
treatment by dermatologists



Clinically Proven Formula



Formulated Without
parabens, oil, phthalates,
dyes and fragrances.



Inspired by
dermatologist
recommended
regimen for
clear skin



Neutrogena Stubborn Acne™ AM Treatment

Vanishing formula reduces size and redness of acne in just hours, with a dermatologist recommended approach.

Use on your full face not just on breakouts. Contains micronized benzoyl peroxide to penetrate deep into pores and kill acne-causing bacteria at the source.

Drug Facts

Active ingredient	Purpose
Benzoyl Peroxide (2.5%)	Acne treatment

Use For the treatment of acne.

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Drug Facts (continued)

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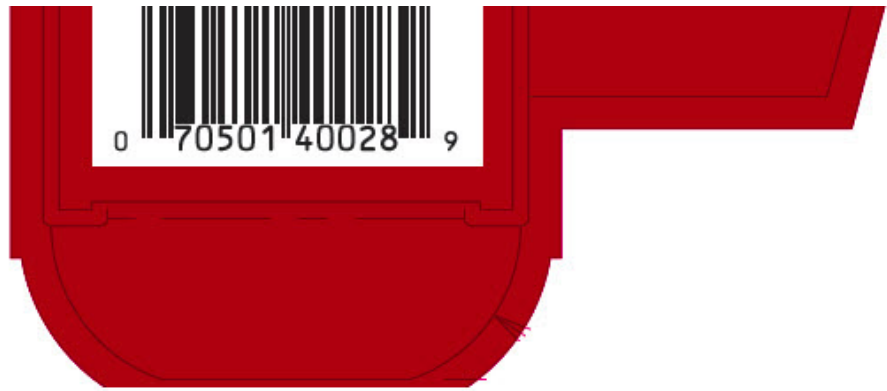
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Made in Canada.

ACTUAL SIZE





NEUTROGENA STUBBORN ACNE AM TREATMENT

benzoyl peroxide gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0653
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	25 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL OR ALLYL SUCROSE CROSSLINKED) (UNII: K6MOM3T5YL)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
LAURETH-4 (UNII: 6HQ855798J)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0653-2	1 in 1 CARTON	08/03/2020	
1		56 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:69968-0653-1	12 in 1 PACKAGE	08/03/2020	

2

9 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	08/03/2020	

Labeler - Kenvue Brands LLC (118772437)

Revised: 11/2024

Kenvue Brands LLC