NEUTROGENA STUBBORN ACNE AM TREATMENT- benzoyl peroxide gel
Johnson & Johnson Consumer Inc.

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.

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

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

STUBBORN ACNE

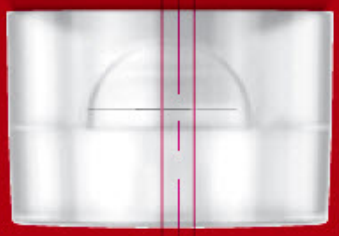
AM TREATMENT

Benzoyl Peroxide
Acne Treatment

works all day to help eliminate stubborn acne

2.5% Micronized BPO

NET WT. 2.0 OZ
(56 g)



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.

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.

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 sunscreen ■ avoid contact with the eyes, lips, and mouth. If contact occurs, flush thoroughly with water. ■ 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.

Do not use if you ■ have very sensitive skin ■ are sensitive to benzoyl peroxide.

Stop use and ask a doctor if ■ irritation becomes severe.

ACTUAL SIZE

Drug Facts (continued)

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

Made in Canada.

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 final	part333D	08/03/2020	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 1/2023

Johnson & Johnson Consumer Inc.