

TARGET UP AND UP ACNE SPOT TREATMENT- benzoyl peroxide lotion
TARGET CORPORATION

Target Up & Up Acne Spot Treatment

Active Ingredient

Benzoyl peroxide 2.5%

Purpose

Acne Medication

Uses

for the treatment of acne

Warnings

For external use only

Do not use if you

- have very sensitive skin
- are sensitive to benzoyl peroxide

When using this product:

- avoid unnecessary sun exposure and use a sunscreen
- 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.
- 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.

Stop use and ask doctor if

- irritation becomes severe.

Keep out of the 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 to 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

- Keep tightly closed
- Avoid storing at extreme temperatures (below 40 F and above 100 F)

Inactive Ingredients

Water, Carbomer, Ethylhexylglycerin, Sodium Hydroxide, Chlorphenesin, Disodium EDTA, Laureth-4, Hydroxypropyl Methylcellulose.

Label

up&up
maximum strength effectiveness
acne spot treatment
benzoyl peroxide acne medication
vanishing formula
NET WT 0.75 OZ (21.2 g)

Drug Facts	
Active ingredient	Purpose
Benzoyl peroxide 2.5%	Acne Medication
Use for the treatment of acne	
Warnings For external use only	
Do not use if you • have very sensitive skin • are sensitive to benzoyl peroxide	
When using this product • avoid unnecessary sun exposure and use a sunscreen • 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. • 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.	
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.	
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up&up maximum strength effectiveness
acne spot treatment
 benzoyl peroxide acne medication
 Compare to Neutrogena® On-the-Spot® Acne Treatment*
 vanishing formula helps reduce acne-causing bacteria won't overdry skin

up&up maximum strength effectiveness
acne spot treatment
 benzoyl peroxide acne medication



NET WT 0.75 OZ (21.2 g)

This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademarks Neutrogena® and Neutrogena On-the-Spot®.



up & up™ maximum strength effectiveness acne spot treatment is fast-acting and helps reduce visible signs of acne blemishes. This vanishing formula penetrates deep to reduce the severity of acne formation of new ones.
 100% satisfaction guaranteed or your money back.
 We welcome any questions you may have at Target.com/ comments or 1-800-810-8874.
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Drug Facts
 Active ingredient: Benzoyl peroxide 2.5% Acne Medication
 Use: For the treatment of acne.
Warnings
 For external use only.
 Do not use if you: have very sensitive skin - are sensitive to benzoyl peroxide.
When using this product
 • avoid unnecessary sun exposure and use a sunscreen
 • avoid contact with the eyes, lips, and mouth.
 • avoid contact with hair and dyed fabrics which may be bleached by this product.
 • skin reactions 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.
 • 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.
 • Benzoyl peroxide is an oxidizing agent. It may bleach or irritate fabrics, including towels, bath linens, and white clothing. It may irritate the eyes.
Keep out of reach of children. If swallowed, get medical help or contact Poison Control Center right away.
Directions
 • Apply the amount shown on the label to the areas affected by acne. Avoid contact with the eyes, lips, and mouth.
 • Apply to dry skin. Do not use on broken skin.
 • Apply to the affected areas of the face and neck. Do not use on the scalp.
Other information
 • Contains benzoyl peroxide.
 • Contains 0.75 oz (21.2 g) of product.
 • Keep this and all medications out of the reach of children.
 • See important information about prescription drugs on the inside of this package.
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TARGET UP AND UP ACNE SPOT TREATMENT

benzoyl peroxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-149
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOYL PEROXIDE (UNII: W9WZ N9A0GM) (BENZOYL PEROXIDE - UNII:W9WZ N9A0GM)	BENZOYL PEROXIDE	25 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
CARBOMER HOMOPOLYMER TYPE B (ALLYL SUCROSE CROSSLINKED) (UNII: Z135WT9208)	
LAURETH-4 (UNII: 6HQ855798J)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-149-51	1 in 1 CARTON	09/13/2016	
1		21 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	09/13/2016	

Labeler - TARGET CORPORATION (006961700)