## NEUTROGENA RAPID CLEAR STUBBORN ACNE SPOT- 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.

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### Neutrogena ® Rapid Clear ® Stubborn Acne Spot Gel

#### **Drug Facts**

#### **Active Ingredients**

Benzoyl Peroxide (10%)

#### **Purpose**

Acne Medication

#### 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.

#### 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 or 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.

#### 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

Protect from Excessive Heat (40 °C/104 °F)

#### **Inactive Ingredients**

water, carbomer homopolymer type B, hydroxypropyl methylcellulose, benzyl alcohol, disodium EDTA, laureth-4, citric acid, sodium hydroxide

#### **Questions?**

Call toll-free **800-582-4048** or **215-273-8755** (collect) or visit www.neutrogena.com Distributed by: **JOHNSON & JOHNSON CONSUMER INC.** Skillman, NJ 08558

#### PRINCIPAL DISPLAY PANEL - 28 g Tube Carton

NFUTROGENA ®

Rapid Clear ® Stubborn Acne Spot Gel

MAXIMUM STRENGTH
BENZOYL PEROXIDE
ACNE MEDICINE
reduces size & redness in just 2 hours

Neutrogena ®

Net Wt. 1 Oz. (28g)

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- ACNE MEDICINE -

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 Cover the entire affected area with a thin layer one to three Purpose Active Ingredient Directions Drug Facts (continued) Drug Facts

Neutrogena® Rapid Clear® Stubborn Acne Spot Gel Clears breakouts with maximum strength Benzoyl Peroxide acne medicine and reduces two key signs of stubborn acne, size and redness, in just two hours.

#### **NEUTROGENA RAPID CLEAR STUBBORN ACNE SPOT**

benzoyl peroxide gel

<b>Product Information</b>	roduct Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0043	
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	100 mg in 1 g	

Inactive Ingredients		
Ingredient Name	Strength	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		
WATER (UNII: 059QF0KO0R)		
CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL OR ALLYL SUCROSE CROSSLINKED) (UNII: K6MOM3T5YL)		

HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
LAURETH-4 (UNII: 6HQ855798J)	

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
NDC:69968- 0043-1	1 in 1 CARTON	09/05/2014		
1	28 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	09/05/2014	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

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