

NEUTROGENA STUBBORN BLACKHEADS DAILY SERUM- salicylic acid lotion
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 BLACKHEADS DAILY SERUM

Drug Facts

Active ingredient

Salicylic Acid 0.5%

Purpose

Acne treatment

Use

- For the treatment of acne
- Penetrates pores to eliminate most blackheads.

Warnings

For external use only.

When using this product

- Avoid contact with eyes. If contact occurs, immediately flush with water.
- 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.

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.

Inactive ingredients

Water, Propylene Glycol, Glycolic Acid, Dimethicone, Mandelic Acid, Gluconolactone, Sodium Hydroxide, Neopentyl Glycol Diheptanoate, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Polyacrylate Crosspolymer-6, Dimethicone Crosspolymer, Disodium EDTA

Other information

- Store at room temperature.
- **Sunburn Alert:** This product contains an alpha hydroxy acid (AHA) that may increase your skin's sensitivity to the sun and particularly the possibility of sunburn. Use a sunscreen, wear protective clothing, and limit sun exposure while using this product and for a week afterwards.

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 - 30 mL Tube Carton

New

Neutrogena®

DERMATOLOGIST

RECOMMENDED BRAND

STUBBORN

BLACKHEADS

DAILY SERUM

Salicylic Acid Acne Treatment

visibly clears clogged pores
and helps eliminate blackheads

10% Glycolic + Mandelic

+ Polyhydroxy Acids

Fragrance-Free

1.0 FL OZ (30mL)



Scan to see your skin's improvement

**Neutrogena
Stubborn Blackheads™ Daily Serum**

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ACTUAL SIZE

Apply 1-2 times daily after cleansing. Use fingertips to gently massage over entire face. This lightweight formula absorbs quickly and is designed to blend effortlessly on all skin tones under moisturizer and/or makeup.

NEW

Neutrogena®

DERMATOLOGIST
RECOMMENDED BRAND



Dermatologist-grade 10% Glycolic + Mandelic + Polyhydroxy Acid Complex is formulated with:

5% Glycolic Acid (AHA) exfoliates to help remove stubborn, pore-clogging debris that can cause acne and make pores look larger

2.5% Mandelic Acid (AHA) Effective exfoliator for oily and acne-prone skin

2.5% Polyhydroxy Acid (PHA) to increase skin clarity without over-drying

Salicylic Acid (BHA) to clear blackheads and help prevent acne from forming



Clinically Proven

Dermatologist tested. Hypoallergenic, non-comedogenic formula. Developed for stubborn acne.



Formulated Without parabens, oil, fragrances, phthalates, dyes

**STUBBORN
BLACKHEADS
DAILY SERUM**

Salicylic Acid Acne Treatment

visibly clears clogged pores
and helps eliminate blackheads

**10% Glycolic + Mandelic
+ Polyhydroxy Acids**

Fragrance-Free



1.0 FL OZ (30 mL)



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NEUTROGENA STUBBORN BLACKHEADS DAILY SERUM

salicylic acid lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0703
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCOLIC ACID (UNII: 0WT12SX38S)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
MANDELIC ACID (UNII: NH496X0UJX)	
GLUCONOLACTONE (UNII: WQ29KQ9POT)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKWBC543X)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (100000 MPA.S AT 1.5%) (UNII: 86FQE96TZ4)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)	
DIMETHICONE CROSSPOLYMER (450000 MPA.S AT 12% IN CYCLOPENTASILOXANE) (UNII: UF7620L1W6)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0703-1	1 in 1 CARTON	08/02/2021	

1		30 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:69968-0703-2	1 in 1 BLISTER PACK	08/02/2021	
2		3.5 mL in 1 POUCH; 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	08/02/2021	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 1/2023

Johnson & Johnson Consumer Inc.