NEUTROGENA CLEAR PORE OIL-ELIMINATING ASTRINGENT- salicylic acid liquid 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 ® Clear Pore Oil-Eliminating Astringent

Drug Facts

Active ingredient

Salicylic Acid 2%

Purpose

Acne medication

Use

For the treatment of acne.

Warnings

For external use only.

Flammable: Keep away from fire or flame.

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 contact with eyes. If contact occurs, flush thoroughly with water.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- cleanse 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.

Other information

Store at Room Temperature.

Inactive ingredients

Water, Alcohol Denat., Hamamelis Virginiana (Witch Hazel) Water, Butylene Glycol, Glycereth-7, Methyl Gluceth-20, Alcohol, Fragrance, Propylene Glycol, Benzophenone-4, Aloe Barbadensis Leaf Extract, Chamomilla Recutita (Matricaria) Flower Extract, Blue 1

Questions?

Call toll-free 800-582-4048 or 215-273-8755 (collect) or visit www.neutrogena.com

Dist. by Neutrogena Corp. Los Angeles CA 90045

PRINCIPAL DISPLAY PANEL - 236 mL Bottle Label

NEW INFO Clear Pore Oil-Eliminating Astringent

effectively treats and helps prevent breakouts without overdrying

salicylic acid acne medication

Neutrogena®

DERMATOLOGIST RECOMMENDED

8 FL. OZ. (236mL)

NEW INFO		Clear skin starts with clear pores. Specifically formulated to instantly remove surface oil to control shine and to treat and help prevent breakouts.
C	lear	Drug Facts Active ingredient Purpose Salicylic Acid 2% Acne medication
	ICOI	Use For the treatment of acne.
P	ore	Warnings For external use only.
Oil-E	-Eliminating Stringent -Eliminating Stringent -Eliminating 	
]	effectively treats and helps prevent breakouts without overdrying	Directions • cleanse 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.
10	salicylic acid	Other information Store at Room Temperature.
Neu	acne medication	Inactive ingredients water, alcohol denat, hamamelis virginiana (witch hazel) water, butylene glycol, glycereth-7, methyl gluceth-20, alcohol, fragrance, propylene glycol, benzophenone-4, aloe barbadensis leaf extract, chamomilla recutita (matricaria) flower extract, blue 1 Questions? Call toll-free 800-582-4048 or 215-273-8755 (collect) or visit www.neutrogena.com
8 FL.	GIST RECOMMENDED 0Z. (236 mL)	Image: The state of the st

NEUTROGENA CLEAR PORE OIL-ELIMINATING ASTRINGENT

salicylic acid liquid

Product Information

Product Type

HUMAN OTC DRUG

TOPICAL

Item Code (Source)

NDC:10812-973

Route of Administration

	t/Active Moiety		
	Ingredient Name	Basis of Strengt	th Strength
Salicylic Acid (UNII: (O414PZ4LPZ) (Salicylic Acid - UNII:O414PZ4LPZ)	Salicylic Acid	20 mg in 1 mL
Inactive Ingredie	nts		
	Strength		
Water (UNII: 059QF01	(O0R)		
Alcohol (UNII: 3K995	3 V9 0 M)		
Hamamelis Virginiaı	na Top Water (UNII: NT00Y05A2V)		
Butylene Glycol (UNI	I: 3XUS85K0RA)		
Glycereth-7 (UNII: 3D	2Y91QZ2H)		
Methyl Gluceth-20 (U	JNII: J3QD0LD11P)		
Propylene Glycol (UN	III: 6DC9Q167V3)		
Sulisobenzone (UNII:	1W6L629B4K)		
Aloe Vera Leaf (UNII	ZY8 1Z8 3H0 X)		
Chamomile (UNII: FG	L3685T2X)		
FD&C Blue No. 1 (UN	II: H3R47K3TBD)		
Packaging			
00	Package Description	Marketing Start Date	Marketing End Dat
# Item Code	Package Description 236 mL in 1 BOTTLE; Type 0: Not a Combination Produc	Marketing Start Date	Marketing End Dat
# Item Code	Package Description 236 mL in 1 BOTTLE; Type 0: Not a Combination Produc	-	Marketing End Da
		-	Marketing End Da
 # Item Code 1 NDC:10812-973-01 	236 mL in 1 BOTTLE; Type 0: Not a Combination Produc	-	Marketing End Da
# Item Code	236 mL in 1 BOTTLE; Type 0: Not a Combination Production Production Production Production	ct 09/15/2011	Marketing End Da Marketing End Da

Labeler - Johnson & Johnson Consumer Inc. (002347102)

Revised: 11/2018

Johnson & Johnson Consumer Inc.