CLEARASIL ULTRA RAPID ACTION SCRUB- salicylic acid lotion
RB Health (US) LLC

Clearasil ®Ultra Rapid Action Scrub

Drug Facts

Active ingredient

Salicylic acid 2%

Purpose

Acne medication

Use

for the treatment of acne

Warnings

For external use only

When using this product

- avoid contact with the eyes. If the product gets into the eyes, rinse thoroughly 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.
- limit use to the face and neck

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

Directions

- wet face
- dispense product into hands and massage gently onto face and neck, avoiding the delicate eye area
- cover the entire affected area with a thin layer and rinse thoroughly with warm water 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 occurs, reduce application to once a day or every other day

Other information

- keep tightly closed
- store in a cool, dry place

Inactive ingredients

water, polyethylene, PPG-15 stearyl ether, glycerin, stearyl alcohol, cetyl betaine, distearyldimonium chloride, sodium lauryl sulfate, cetyl alcohol, alcohol, steareth-21, sodium chloride, behenyl alcohol, synthetic wax, steareth-2, fragrance, Lavandula stoechas extract, Helichrysum italicum extract, Cistus monspeliensis extract, xanthan gum, dimethyl palmitamine, lauryl alcohol, mica, isopropyl alcohol, disodium EDTA, BHT, magnesium nitrate, methylchloroisothiazolinone, magnesium chloride, methylisothiazolinone, sodium sulfate, ferric ferrocyanide, titanium dioxide, FD&C blue no. 1

Questions?

call **1-866-25-CLEAR (1-866-252-5327).**
You may also report side effects to this phone number

Distributed by: Reckitt Benckiser LLC
Parsippany, NJ 07054-0224

Made in France

PRINCIPAL DISPLAY PANEL - 150 mL Tube Label

Clearasil ®

ULTRA

Rapid Action
Face Scrub

Salicylic Acid 2% Acne Medication

MAXIMUM STRENGTH
12
HRS
Visibly
clearer skin
in as fast as
12 hours

5 FL OZ (150 mL)

CLEARASIL ULTRA RAPID ACTION SCRUB

salicylic acid lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63824-416
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	2 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	
PPG-15 STEARYL ETHER (UNII: 1II18XLS1L)	
GLYCERIN (UNII: PDC6A3C0OX)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	

CETYL BETAINE (UNII: E945X08YA9)	
DISTEARYLDIMONIUM CHLORIDE (UNII: OM9573ZX3X)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ALCOHOL (UNII: 3K9958V90M)	
STEARETH-21 (UNII: 53J3F32P58)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
DOCOSANOL (UNII: 9G1OE216XY)	
STEARETH-2 (UNII: V56DFE46J5)	
LAVANDULA STOECHAS FLOWERING TOP (UNII: 70759G2U6A)	
HELICHRYSUM ITALICUM FLOWER (UNII: P62Y550X24)	
XANTHAN GUM (UNII: TTV12P4NEE)	
DIMETHYL PALMITAMINE (UNII: 5E4QI660PW)	
LAURYL ALCOHOL (UNII: 178A96NLP2)	
MICA (UNII: V8A1AW0880)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
FERRIC FERROCYANIDE (UNII: TLE294X33A)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63824-416-65	150 mL in 1 TUBE; Type 0: Not a Combination Product	12/01/2013	09/01/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	12/01/2013	09/01/2024

Labeler - RB Health (US) LLC (081049410)

Revised: 10/2023 RB Health (US) LLC