

CLEARASIL ULTRA RAPID ACTION TREATMENT- salicylic acid lotion
Reckitt Benckiser LLC

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.

Clearasil® Ultra
Rapid Action Treatment Lotion

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 product gets into the eyes rinse thoroughly with water.
- using other topical acne medications at the same time or immediately following the use of this product, may increase dryness or irritation of the skin. If this occurs, only one acne medication should be used unless directed by your doctor.
- limit use to the face and neck, chest and back

Stop use and ask a doctor if skin or eye irritation develops

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

Directions

- cleanse 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 2 or 3 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 in a cool, dry place

Inactive ingredients

water, butylene glycol, octyldodecanol, steareth-2, cetyl alcohol, steareth-21, dimethicone,

polyacrylamide, glycerin, C13-14 isoparaffin, sodium hydroxide, xanthan gum, magnesium aluminum silicate, laureth-7, fragrance, Lavandula stoechas extract, Helichrysum italicum extract, Cistus monspeliensis extract, titanium dioxide

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

PRINCIPAL DISPLAY PANEL - 118 mL Tube Label

NEW

Clearasil®

ULTRA

Rapid Action

Treatment Lotion

Face, Chest & Back

Salicylic Acid 2% Acne Medication

4

HRS

MAXIMUM STRENGTH

Visibly reduces

pimple size in

as fast as

4 hours

With Acceladerm® Technology

4 FL. OZ. (118 mL)

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Face, Chest & Back
Salicylic Acid 2% Acne Medication

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MAXIMUM STRENGTH

Visibly reduces
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With Acceladerm® Technology

4 FL. OZ. (118 mL)

When treating breakouts, faster is better. Clearasil Ultra™ Rapid Action Treatment Lotion works with the skin to unblock pores and deliver max strength active ingredient into the pores to treat pimples - in as fast as 4 hours!

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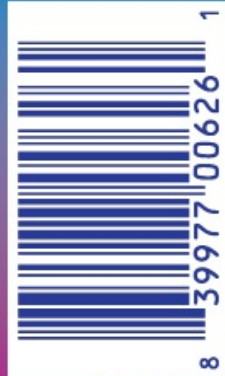
water, butylene glycol, octyldodecanol, steareth-2, cetyl alcohol, steareth-21, dimethicone, polyacrylamide, glycerin, C13-14 isoparaffin, sodium hydroxide, xanthan gum, magnesium aluminum silicate, laureth-7, fragrance, Lavandula stoechas extract, Helichrysum italicum extract, Cistus monspeliensis extract, titanium dioxide

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salicylic acid lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63824-314
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: 059QF0K00R)	
butylene glycol (UNII: 3XUS85K0RA)	
octyldodecanol (UNII: 461N1O614Y)	
cetyl alcohol (UNII: 936JST6JCN)	
steareth-21 (UNII: 53J3F32P58)	
dimethicone (UNII: 92RU3N3Y1O)	
glycerin (UNII: PDC6A3C0OX)	
C13-14 isoparaffin (UNII: E4F12ROE70)	
sodium hydroxide (UNII: 55X04QC32I)	
xanthan gum (UNII: TTV12P4NEE)	
magnesium aluminum silicate (UNII: 6M3P64V0NC)	
laureth-7 (UNII: Z95S6G8201)	
Helichrysum Italicum Flower (UNII: P62Y550X24)	
titanium dioxide (UNII: 15FIX9V2JP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63824-314-01	118 mL in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part333D	11/08/2013	

Labeler - Reckitt Benckiser LLC (094405024)

Revised: 11/2013

Reckitt Benckiser LLC