

CLEARASIL ULTRA ACNE PLUS MARKS DAILY SCRUB- 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 Acne Plus Marks Daily 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 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 at 20-25°C (68-77°F)

Inactive ingredients

water, sodium cocoyl isethionate, glycerin, cetyl alcohol, laureth-3, polyethylene, sodium laureth sulfate, sodium cocoamphoacetate, niacinamide, sodium lactate, sorbitol, hexyldecanol, fragrance, acrylates/C10-30 alkyl acrylate crosspolymer, sodium hydroxide, dipotassium glycyrrhizate, disodium EDTA, bisabolol, cetylhydroxyproline palmitamide, stearic acid, polyquaternium-43, Brassica campestris (rapeseed) sterols, phenoxyethanol, sodium benzoate, FD&C blue no. 1, FD&C yellow no. 6

Questions?

call **1-866-25-CLEAR (1-866-252-5327)**.

You may also report side effects to this phone number

Distributed by:

Reckitt Benckiser

Parsippany, NJ 07054-0224

Made in USA

PRINCIPAL DISPLAY PANEL - 150 mL Tube Label

Clearasil®

ULTRA

Acne + Marks

Daily Scrub

Salicylic Acid 2% Acne Medication

REDUCES

MARKS

**Helps clear
pimples fast,
and reduces the
appearance of
acne marks**

5.0 FL OZ (150 mL)

Clearasil®

ULTRA

Acne + Marks Daily Scrub

Salicylic Acid 2% Acne Medication

Helps clear
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and reduces the
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5.0 FL OZ (150 mL)

Clearasil ULTRA® Skin science says:
Acne marks actually start to form when a pimple appears and can take weeks to go away.
How Clearasil® works with your skin
With dermatologist-inspired science, this Scrub is designed to work on your skin from the start of the pimple. It contains a unique ingredient complex, with Vitamin B3 known to reduce the appearance of marks and exfoliates to enhance the skin renewal process. The scrubbing beads provide a deep cleansing experience.
Also available, Clearasil ULTRA® Acne+Marks Wash & Mask.

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www.clearasil.us

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Reckitt Benckiser
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CLEARASIL ULTRA ACNE PLUS MARKS DAILY SCRUB

salicylic acid lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63824-420
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)
SODIUM COCOYL ISETHIONATE (UNII: 518XTE8493)
GLYCERIN (UNII: PDC6A3C0OX)
CETYL ALCOHOL (UNII: 936JST6JCN)
LAURETH-3 (UNII: F32E4CB0UJ)
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)
SODIUM COCOAMPHOACETATE (UNII: W7Q5E87674)
NIACINAMIDE (UNII: 25X51I8RD4)
SODIUM LACTATE (UNII: TU7HW0W0QT)
SORBITOL (UNII: 506T60A25R)
HEXYLDECANOL (UNII: 151Z7P1317)
SODIUM HYDROXIDE (UNII: 55X04QC32I)
GLYCYRRHIZINATE DIPO TASSIUM (UNII: CA2Y0FE3FX)
EDETATE DISODIUM (UNII: 7FLD91C86K)
LEVOMENOL (UNII: 24WE03BX2T)
CETYLHYDROXYPROLINE PALMITAMIDE (UNII: 74ONU0S62G)
STEARIC ACID (UNII: 4ELV7Z65AP)
RAPESEED STEROL (UNII: B46B6DD20U)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
SODIUM BENZOATE (UNII: OJ245FE5EU)
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63824-420-65	150 mL in 1 TUBE; Type 0: Not a Combination Product	12/22/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part333D	12/22/2014	

Labeler - Reckitt Benckiser LLC (094405024)