UP AND UP DUAL EFFECTS MOISTURIZER- salicylic acid lotion TARGET CORPORATION

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.

Up & Up Dual Effects Moisturizer

Active ingredient

Salicylic Acid 0.5%

Purpose

Acne Medication

Uses

• For the management of acne.

Warnings

For external use only

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 doctor. • If bothersome dryness or peeling occurs, reduce application to once a day or every other day.

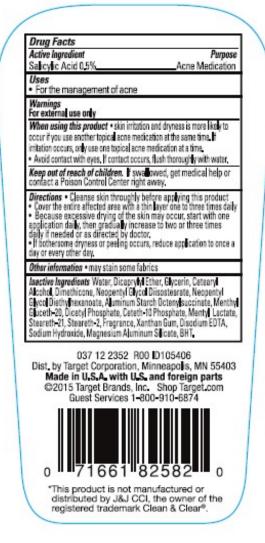
Other information

• may stain some fabrics

Inactive ingredients

Water, Dicaprylyl Ether, Glycerin, Cetearyl Alcohol, Dimethicone, Neopentyl Glycol Diisostearate, Neopentyl Glycol Diethylhexanoate, Aluminum Starch Octenylsuccinate, Methyl Gluceth-20, Dicetyl Phosphate, Ceteth-10 Phosphate, Mentyl Lacatate, Steareth-21, Steareth-2, Fragrance, Xanthan Gum, Disodium EDTA, Sodium Hydroxide, Magnesium Aluminum Silicate, BHT.





UP AND UP DUAL EFFECTS MOISTURIZER

salicylic acid lotion

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-931
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)	SALICYLIC ACID	5 mg in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
STEARETH-2 (UNII: V56DFE46J5)		
GLYCERIN (UNII: PDC6A3C0OX)		
DICAPRYLYL ETHER (UNII: 77JZM5516Z)		
WATER (UNII: 059QF0KO0R)		

STEARETH-21 (UNII: 53J3F32P58)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: 19PJ0O6294)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
DIMETHICONE (UNII: 92RU3N3Y10)	
NEOPENTYL GLYCOL DIISOSTEARATE (UNII: 4M6OQ34JWW)	
CETETH-10 PHOSPHATE (UNII: 4E05O5N49G)	
NEOPENTYL GLYCOL DIETHYLHEXANOATE (UNII: U68ZV6W62C)	
MENTHYL LACTATE, (-)- (UNII: 2BF9E65L7I)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
XANTHAN GUM (UNII: TTV12P4NEE)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
DIHEXADECYL PHOSPHATE (UNII: 2V6E5WN99N)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673- 931-53	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/11/2012	

Marketing In	larketing Information		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	10/11/2012	

Labeler - TARGET CORPORATION (006961700)

Revised: 10/2022 TARGET CORPORATION