UP AND UP ILLUMINATING DAILY MOISTURIZER WITH SUNSCREEN BROAD SPECTRUM SPF 15- avobenzone, octinoxate, octisalate 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 and Up Illuminating Daily Moisturizer with Sunscreen Broad Spectrum SPF 15

Active ingredients

Avobenzone 3.0%, Octinoxate 7.5%, Octisalate 2.0%

Purpose

Sunscreen

Uses

- helps prevent sunburn
- if used as directed with other sun protection measures (see ***Directions***), decreases the risk of skin cancer and early aging caused by the sun

Warnings

For external use only

Do not use

• on damaged or broken skin.

When using this product

• keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if

• rash occurs.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply liberally 15 minutes before sun exposure.
- reapply:
- after 80 minutes of swimming or sweating
- immediately after towel drying
- at least every 2 hours
- children under 6 months of age: Ask a doctor
- **Sun Protection Measures**. Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a Broad Spectrum SPF value of 15 or higher and other sun protection measures including:
- limit time in the sun especially from 10 a.m. – 2 p.m.
- wear long-sleeved shirts, pants, hats, and sunglasses

Other information

- protect the product in this container from excessive heat and direct sun
- may stain or damage some fabrics, materials or surfaces

Inactive ingredients

Water, C12-15 Alkyl Benzoate, Glycerin, Cetearyl Alcohol, Dimethicone, Glycine Soja (Soybean) Seed Extract, Phenyl Trimethcione, Arachidyl Alcohol, Cetearyl Glucoside, Phenoxyethanol, Benzyl Alcohol, Panthenol, Ethylene/ Acrylic Acid Copolymer, Behenyl Alcohol, Steareth-2, Fragrance, Steareth-21, Polymethyl Methacrylate, Polyacrylamide, Arachidyl Glucoside, Disodium EDTA, C13-14 Isoparaffin, Laureth-7, Silica, Benzalkonium Chloride, Iodopropynyl Butylcarbamate, PEG-4 Dilaurate, PEG-4 Laurate, PEG-4, Alcohol, BHT, Sodium Hydroxide, Citric Acid, Titanium Dioxide, Mica

Up and Up Illuminating Daily Moisturizer with Sunscreen Broad Spectrum SPF 15

4 FL OZ (118.3 mL)

NDC 11673-934-04 Bottle

NDC 11673-934-05 Carton

UP AND UP ILLUMINATING DAILY MOISTURIZER WITH SUNSCREEN BROAD SPECTRUM SPF 15

avobenzone, octinoxate, octisalate lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-934
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	20 mg in 1 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
GLYCOL DILAURATE (UNII: 9O691KKR2A)	
PEG-4 LAURATE (UNII: AYF4VM3N1Z)	

BENZYL ALCOHOL (UNII: LKG8494WBH)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
ACRYLIC ACID/ETHYLENE COPOLYMER (600 MPA.S) (UNII: 1PEZ3NLY6I)	
POLYACRYLAMIDE (1500 MW) (UNII: 5D6TC4BRWV)	
ARACHIDYL GLUCOSIDE (UNII: 6JVW35JOOJ)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SOYBEAN OIL (UNII: 241ATL177A)	
PANTHENOL (UNII: WV9CM0O67Z)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
STEARETH-21 (UNII: 53J3F32P58)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
ARACHIDYL ALCOHOL (UNII: 1QR1QRA9BU)	
PHENYL TRIMETHICONE (UNII: DR0K5NOJ4R)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
POLY(METHYL METHACRYLATE; 450000 MW) (UNII: Z47NNT4J11)	
POLYETHYLENE GLYCOL 200 (UNII: R95B8J264J)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
WATER (UNII: 059QF0KO0R)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)	
DOCOSANOL (UNII: 9G1OE216XY)	
MICA (UNII: V8A1AW0880)	
LAURETH-7 (UNII: Z95S6G8201)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
STEARETH-2 (UNII: V56DFE46J5)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-934-05	1 in 1 CARTON	02/01/2013	
1	NDC:11673-934-04	118.3 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	02/01/2013	

Labeler - TARGET CORPORATION (006961700)