

TOUCH ANTI-AGING SPF 30 MOISTURIZER- zinc oxide, octinoxate, and octisalate cream

Diractions Capital 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.

Touch™ Anti-Aging SPF 30 Moisturizer

Drug Facts

ACTIVE INGREDIENTS

Zinc Oxide 7.0%, Octinoxate 7.5%, Octisalate 5.0%

PURPOSE

sunscreen

Uses

helps prevent sunburn.

Warnings

For external use only

- avoid contact with eyes
- discontinue use if signs of irritation or rash appear
- **Stop use and consult a doctor** if irritation or rash persists.

Directions

Apply to areas exposed to sun. Reapply frequently, especially after physical activity or swimming.

Sun Alert

Limiting sun exposure, wearing protective clothing, and using sunscreens may reduce the risks of skin aging, skin cancer, and other harmful effects of the sun.

Inactive Ingredients

Aqua, C12-15 Alkyl Benzoate, Glyceryl Stearate, Cetyl Alcohol, Cetearyl Ethylhexanoate, Cyclopentasiloxane, Cetearyl Alcohol, Cetyl Dimethicone, Ascorbic Acid, Tocopherol Acetate, Calendula Officinalis Flower Extract, Chamomilla Recutita (Matricaria) Flower Extract, Ginkgo Biloba Leaf Extract, PEG-100 Stearate, Panax Ginseng Root Extract,

Camellia Oleifera Leaf Extract, Glycyrrhiza Glabra (Licorice) Root Extract, Sodium PCA, Sodium Hyaluronate, Xanthan Gum, Dehydroacetic Acid, Triethoxycaprylylsilane, Benzyl Alcohol, Tetrasodium EDTA, Citric Acid.

Distributed by
Touch Skin Care
Bellaire, TX 77401

PRINCIPAL DISPLAY PANEL - 100 mL Bottle Label

touch™

anti-aging
SPF 30
moisturizer

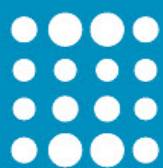
Ultra-Light Broad Spectrum UVA/UVB
with Hyaluronic Acid & Vitamin C & E
Oil-Free & Fragrance-Free

cruelty free
vegan

lotion
SPF
MADE IN
USA

Shake Well
Before Use

3.3 FL. OZ.
(100mL)



touch™

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Touch Skin Care
Bellaire, TX 77401
MADE IN
USA



X002HF43G9

Touch Anti-Aging SPF 30...NEW

MyTouchSkincare.com

TOUCH ANTI-AGING SPF 30 MOISTURIZER

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73246-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Zinc Oxide (UNII: SOI2LOH54Z) (Zinc Oxide - UNII:SOI2LOH54Z)	Zinc Oxide	70 mg in 1 mL
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	75 mg in 1 mL
Octisalate (UNII: 4X49Y0596W) (Octisalate - UNII:4X49Y0596W)	Octisalate	50 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	

ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CETEARYL ETHYLHEXANOATE (UNII: 9M64UO4C25)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETYL DIMETHICONE 45 (UNII: IK315POC44)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
MATRICARIA CHAMOMILLA FLOWERING TOP (UNII: 3VNC7T6Z02)	
GINKGO BILOBA WHOLE (UNII: 660486U6OI)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PANAX GINSENG ROOT OIL (UNII: P9T4K47OM0)	
CAMELLIA OLEIFERA LEAF (UNII: 5077EL0C60)	
GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
XANTHAN GUM (UNII: TTV12P4NEE)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
EDETATE SODIUM (UNII: MP1J8420LU)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73246-002-01	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/04/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part352	03/04/2020	

Labeler - Diractions Capital LLC (117719327)