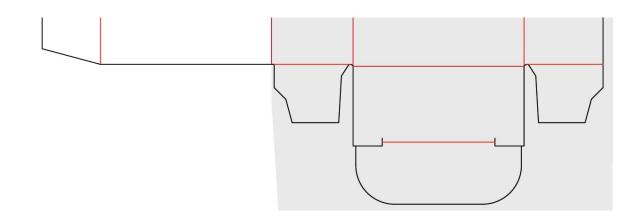
# POST RAYS UV PROTECT SUN BLOCK- zinc oxide, octinoxate, octisalate, titamium dioxide cream Humex, Inc

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.

-----

# Drug Facts Active Ingredient Purpose





Zinc Oxide (5.006%)	Sunscreen
Octinoxate (6.8%)	Sunscreen
Octisalate (4%)	Sunscreen
Titanium Dioxide (4.0014%)	Sunscreen

#### Uses

Helps prevent sunburn

## **Warnings**

**Skin Cancer/Skin Aging Aler**t: Spending time in the sun increases your risk of skin cancer and early skin aging. This product has been shown only to help prevent sunburn, not skin cancer or early skin aging.

## 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 (1-800-222-1222) right away.

#### Directions

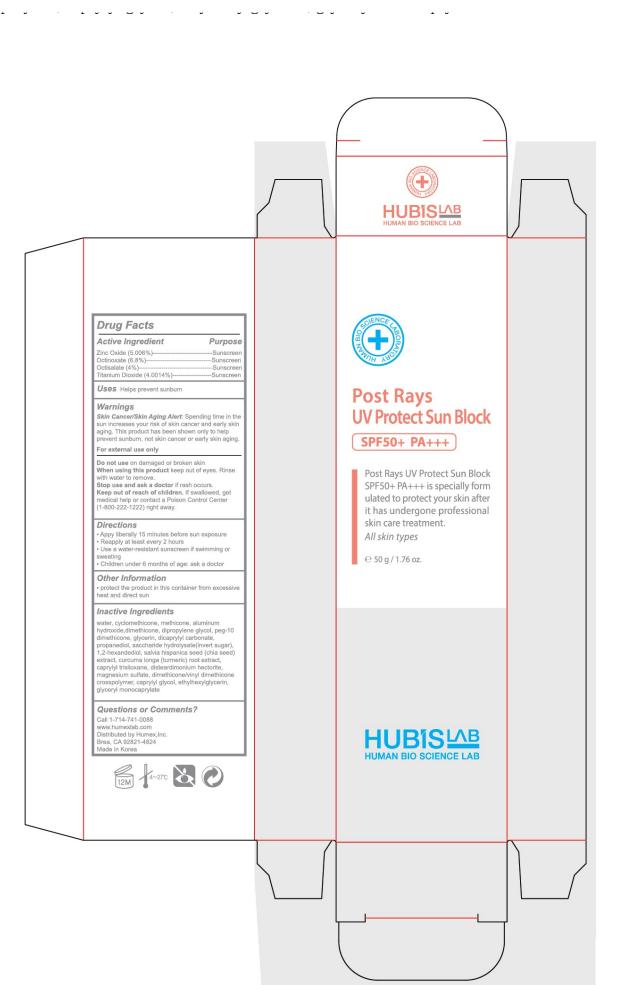
- Apply liberally 15 minutes before sun exposure
- Reapply at least every 2 hours
- Use a water-resistant sunscreen if swimming or sweating
- Children under 6 months of age: ask a doctor

### Other Information

protect the product in this container from excessive heat and direct sun

# **Inactive Ingredients**

water, cyclomethicone, methicone, aluminum hydroxide, dimethicone, dipropylene glycol, peg-10 dimethicone, glycerin, dicaprylyl carbonate, propanediol, saccharide hydrolysate(invert sugar), 1,2-hexandediol, salvia hispanica seed (chia seed) extract, curcuma longa (turmeric) root extract, caprylyl trisiloxane, disteardimonium hectorite, magnesium sulfate, dimethicone/vinyl dimethicone crosspolymer, caprylyl glycol, ethylhexylglycerin, glyceryl monocaprylate



# POST RAYS UV PROTECT SUN BLOCK

zinc oxide,octinoxate, octisalate,titamium dioxide cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76344-8002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	6.8 g in 100 g	
OCTISALATE (UNII: 4X49 Y0596W) (OCTISALATE - UNII:4X49 Y0596W)	OCTISALATE	4.5 g in 100 g	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP) (TITANIUM DIO XIDE - UNII:15FIX9 V2JP)	TITANIUM DIO XIDE	4.0014 g in 100 g	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	5.006 g in 100 g	

Inactive Ingredients				
Ingredient Name	Strength			
CAPRYLYL TRISILO XANE (UNII: Q95M2P1KJL)	1 g in 100 g			
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	0.8 g in 100 g			
DISTEARDIMO NIUM HECTO RITE (UNII: X687XDK09L)	0.8 g in 100 g			
TURMERIC (UNII: 856 YO 1Z64F)	0.01 g in 100 g			
GLYCERYL CAPRYLATE (UNII: TM2TZD4G4A)	0.1 g in 100 g			
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	3 g in 100 g			
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	0.1 g in 100 g			
CYCLOMETHICONE 5 (UNII: 0 THT5PCI0 R)	13 g in 100 g			
DIPROPYLENE GLYCOL (UNII: E107L85C40)	3 g in 100 g			
INVERT SUGAR (UNII: ED959S6ACY)	0.1 g in 100 g			
SALVIA HISPANICA SEED (UNII: NU0 OLX0 6 F8)	0.01 g in 100 g			
CAPRYLYL GLYCOL (UNII: 00 YIU5438 U)	0.2 g in 100 g			
ALUMINUM HYDRO XIDE (UNII: 5QB0T2IUN0)	0.5643 g in 100 g			
PROPANEDIOL (UNII: 5965N8W85T)	0.1 g in 100 g			
1,2-HEXANEDIOL (UNII: TR046 Y3K1G)	0.02 g in 100 g			
DICAPRYLYL CARBONATE (UNII: 609 A3V1SUA)	2 g in 100 g			
METHICONE (20 CST) (UNII: 6777U11MKT)	0.2084 g in 100 g			
DIMETHICO NE (UNII: 92RU3N3Y1O)	0.2565 g in 100 g			
GLYCERIN (UNII: PDC6A3C0OX)	2 g in 100 g			

Product Characteristics			
Color	white (Milky White)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date
1	NDC:76344-8002-2	1 in 1 BOX	12/0 1/20 19	
1	NDC:76344-8002-1	50 g in 1 TUBE; 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	12/0 1/20 19	

# Labeler - Humex,Inc (017945311)

Establishment			
Name	Address	ID/FEI	Business Operations
Kolmar Korea		689512611	manufacture(76344-8002)

Revised: 1/2021 Humex,Inc