

PURPOSE- octinoxate, octisalate and oxybenzone lotion
Bausch & Lomb Incorporated

Drug Facts

Active ingredients

Octinoxate 7.5%
Octisalate 5%
Oxybenzone 3%

Purpose

Sunscreen

Use

■ helps prevent sunburn.

Warnings

Skin Cancer/Skin Aging Alert: 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 product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply generously and evenly 15 minutes before sun exposure
- apply to all skin exposed to the sun
- use a water resistant sunscreen if

- swimming or sweating
- reapply at least every 2 hours
- children under 6 months of age:
Ask a doctor

Other information

- protect this product from excessive heat and direct sun
- may stain or damage some fabrics or surfaces

Inactive ingredients

water, octyldodecyl neopentanoate, glycerin, emulsifying wax NF, glyceryl stearate, PEG-100 stearate, dimethicone, triethanolamine, diazolidinyl urea, carbomer, methylparaben, ethylparaben, propylparaben, bisabolol, farnesol, citric acid

Questions or comments?

Call 1-800-553-5340

Principal Display Panel

PURPOSE®

Dual
Treatment
Moisture
Lotion

Sunscreen SPF 10

Oil-Free

Fragrance Free

Provides
Long-Lasting
Moisture

Developed with
Dermatologists

4 OZ (113g)



PURPOSE

octinoxate, octisalate and oxybenzone lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:24208-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7.5 g in 100 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 g
OXYBENZONE (UNII: 950OS7VE0Y) (OXYBENZONE - UNII:950OS7VE0Y)	OXYBENZONE	3 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
TROLAMINE (UNII: 9O3K93S3TK)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
ETHYLPARABEN (UNII: 14255EXE39)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
LEVOMENOL (UNII: 24WE03BX2T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24208-001-04	1 in 1 CARTON	10/07/2013	
1		113 g 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 Drug	M018	10/07/2013	

Labeler - Bausch & Lomb Incorporated (196603781)

Establishment

Name	Address	ID/FEI	Business Operations
Voyant Beauty, Inc.		243547333	manufacture(24208-001)

Revised: 9/2024

Bausch & Lomb Incorporated