

KROGER SUNSCREEN SPF 8- octinoxate oxybenzone lotion
THE KROGER CO

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 ingredients

Octinoxate 7.5%

Oxybenzone 2.0%

Purpose

Sunscreen

Uses

- helps prevent sunburn
- higher SPF gives more sunburn protection
- retains SPF after 80 minutes of activity in the water

Warnings

For external use only

When using this product

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

stop use and ask a doctor if

- rash or irritation develops and lasts.

keep out of reach of children.

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

Directions

- apply generously and evenly before sun exposure and as needed
- children under 6 months of age:ask a doctor
- reapply frequently and after towel drying, swimming or perspiring.

Other Information

- May stain some fabrics
- 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

Water, Ethylhexyl Palmitate, Propylene Glycol, Cetyl Phosphate, Triethanolamine, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Carbomer, Tocopherol, Disodium EDTA, Aloe Barbadensis Leaf Juice Powder, Diazolidinyl Urea, Methylparaben, Propylparaben, Fragrance.

For comments or questions
please call 1-800-632-6900

Principal Display Panel

croger

QUALITY GUARANTEED

Sunscreen 8 spf

SUNSCREEN LOTION

with

Aloe and Vitamin E

Very

Water

Resistant

8 FL OZ (236 mL)



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Distributed by
The Kroger Co.,
Cincinnati, Ohio 45202

Lotion Made in the U.S.A.
Packaging Made in Canada

KR009-A

KROGER SUNSCREEN SPF 8

octinoxate oxybenzone lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:30 142-925
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
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CETYL PHOSPHATE (UNII: VT07D6X67O)	
TROLAMINE (UNII: 9O3K93S3TK)	
TOCOPHEROL (UNII: R0ZB2556P8)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30142-925-56	226 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	11/12/2012	

Labeler - THE KROGER CO (006999528)

Revised: 11/2012

THE KROGER CO