RADIABLOCK- oxybenzone, octinoxate lipstick Medline Industries, LP

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.

094 RadiaBlock Sunblock & LipBalm

Active ingredient

Oxybenzone 3%

Padimate O 7%

Purpose

Sunscreen

Sunscreen

Uses

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 broken or damaged skin

Stop use and ask a doctor

if rash occurs

When using this product

keep out of eyes. Rinse with water to remove.

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

Use a water resistant sunscreen if swimming or sweating

Reaaply at least every 2 hours

Children under 6 months: Ask a doctor

Inactive ingredients

Aloe Barbadensis (Aloe Vera) Leaf Extract, Copernicia Cerifera (Carnauba) Wax, Flavor, Mineral Oil, Paraffin, Petrolatum, Retinyl Palmitate, Simmondsia Chinensis (Jojoba) Seed Oil, Theobroma Cacao (Cocoa) Seed Butter, Tocopherol (Vitamin E), Zea Mays (Corn) Oil (and) BHA (and) Propyl Gallate (and) Citric Acid

Other information

- protect the product in this container from excessive heat and direct sun
- do not use if safety seal is broken or missing

Questions or comments?

1-800-MEDLINE (633-5463)

Monday-Friday, 9AM-5PM CST

Principal Display Panel

SPF 15 Lip Balm

NDC 53329-094-83

Carrington

RadiaBlock

Sunscreen SPF 15

Net Wt: .015 oz/4.2g



RADIABLOCK

oxybenzone, octinoxate lipstick

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:53329-094 Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OXYBENZONE (UNII: 9500S7VE0Y) (OXYBENZONE - UNII:9500S7VE0Y)	OXYBENZONE	30 mg in 1 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 g

Inactive Ingredients		
Ingredient Name	Strength	
COCOA BUTTER (UNII: 5120YT1CRR)		
.ALPHATOCOPHEROL (UNII: H4N855PNZ1)		
CORN OIL (UNII: 8470G57WFM)		
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
CARNAUBA WAX (UNII: R12CBM0EIZ)		
MINERAL OIL (UNII: T5L8T28FGP)		
PARAFFIN (UNII: 1900E3H2ZE)		
PETROLATUM (UNII: 4T6H12BN9U)		
PROPYL GALLATE (UNII: 8D4SNN7V92)		
JOJOBA OIL (UNII: 724GKU717M)		

l	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:53329-094- 83	4.2 g in 1 TUBE; Type 0: Not a Combination Product	03/01/2013	04/30/2025	

Marketing In	keting Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part352	03/01/2013	04/30/2025		

Labeler - Medline Industries, LP (025460908)

Registrant - Medline Industries, LP (025460908)

Revised: 8/2023 Medline Industries, LP