CHAPICE SPF 30 LIP BALM- octinoxate, octocrylene, oxybenzone, octisalate stick
OraLabs

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

Octinoxate (7.5%), Octocrylene (10.0%), Oxybenzone (5.0%), Octisalate (5.0%)

Purpose

Sunscreen, Sunscreen, Sunscreen, Sunscreen

Keep Out of Reach of Children

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

Uses

Prevents Sunburns

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 prevent sunburn, not skin cancer or early aging.

For external use only: Stop use and ask a doctor: if rash or irritation develops and lasts.

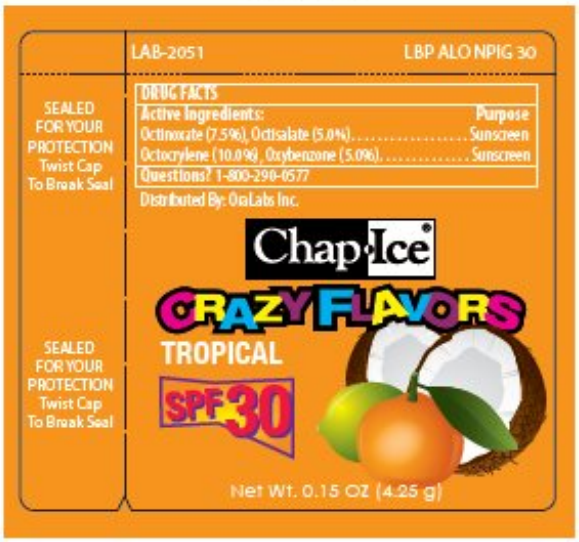
Directions

Apply liberally before sun exposure and as needed. Children under 6 months of age: Ask a doctor before use.

Inactive Ingredients

Mineral Oil, Ozokerite, Petrolatum, Microcrystalline Wax, Ceresin, Flavor, Copernicia Cerifera (Carnauba) Wax, Caprylic/Capric Triglyceride, Titanium Dioxide (CI 77891), Stearic Acid, Tocopheryl Acetate, Aluminum Hydroxide.

Package/Label Principal Display Panel



CHAPICE SPF 30 LIP BALM

octinoxate, octocrylene, oxybenzone, octisalate stick

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63645-164
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7.5 mg in 1 g
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	5.0 mg in 1 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	10.0 mg in 1 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5.0 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	13.0 mg in 1 g
LIGHT MINERAL OIL (UNII: N6K5787QVP)	25.0 mg in 1 g
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	.5 mg in 1 g
CERESIN (UNII: Q1LS2UJO3A)	27 mg in 1 g
MICROCRYSTALLINE WAX (UNII: XOF597Q3KY)	4 mg in 1 g

Product Characteristics

Color	WHITE	**Score**	
Shape		**Size**	
Flavor	COCONUT	**Imprint Code**	

Contains	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63645-164-01	1 g in 1 CONTAINER; Type 0: Not a Combination Product	10/09/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	10/09/2017	

Labeler - OraLabs (801824756)

Registrant - OraLabs (801824756)

Establishment

Name	Address	ID/FEI	Business Operations
OraLabs		801824756	MANUFACTURE(63645-164) , LABEL(63645-164)

Revised: 11/2022 OraLabs