PEARLIXIME ILLUMINATING PERFECTING SUNSCREEN BROAD SPECTRUM SPF 15-avobenzone, octinoxate, octocrylene cream
IXXI S.A.S.

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.

Pearlixime ILLUMINATING PERFECTING CREAM Sunscreen Broad Spectrum SPF 15

Drug Facts

Active ingredients

Avobenzone 3%

Octinoxate 7.5%

Octocrylene 10%

Purpose

Sunscreen

Uses

- Helps prevent sunburn.
- If used as directed with other sun protection measures (see ***Directions***), decreases the risk of skin cancer and early skin aging caused by the sun.

Warnings

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 ues and ask a doctor if

- rash occurs

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
• Reapply at least every 2 hours

•Use a Water Resistant sunscreen if swimming or sweating

• **Sun Protection Measures.** Speanding time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a broad spectrum SPF value of 15 or higher and other sun protection measures including:

• Limit time in the sun, especially from 10 a.m. - 2 p.m.
• Wear long-sleeved shirts, pants, hats, and sunglasses
• Children under 6 months of age: Ask a doctor.

Other Information

• Protect the product inthis container from excessive heat and direct sun.

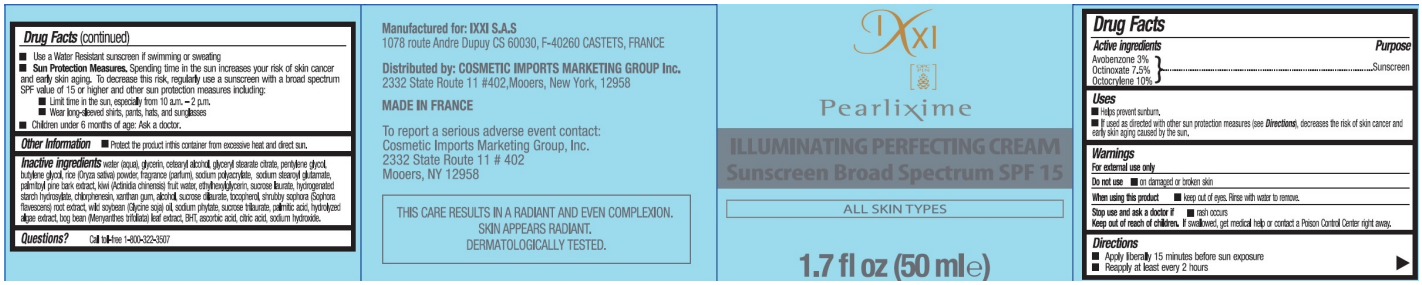
Inactive ingredients

water (aqua), glycerin, cetearyl alcohol, glyceryl stearate citrate, pentylene glycol, butylene glycol, rice (Oryza sative) powder, fragrance (parfum), sodium polyacrylate, sodium stearoyl glutamate, palmitoyl pine bark extract, kiwi (Actinidia chinensis) fruit water, ethylhexylglycerin, sucrose laurate, hydrogenated starch hydrosylate, chlorphenesin, xanthan gum, alcohol, sucrose dilaurate, tocopherol, shrubby sophota (sophora flavescens) root extract, wild soybean (glycine soja) oil, sodium phytate, sucrose trilaurate, palmitic acid, hydrolyzed algae extract, bog bean (Menyanthes trifoliata) leaf extract, BHT, ascorbic acid, citric acid, sodium hydroxide.

Questions?

Call toll-free 1-800-322-3507

Package Labeling:



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avobenzone, octinoxate, octocrylene cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70532-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	100 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
GLYCERYL STEARATE CITRATE (UNII: WH8T92A065)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
RICE BRAN (UNII: R60QEP13IC)	
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)	
KIWI FRUIT OIL (UNII: 66086CWP3Q)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
SUCROSE LAURATE (UNII: 05Q7CD0E49)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ALCOHOL (UNII: 3K9958V90M)	
SUCROSE DILAURATE (UNII: 5926LC4S7M)	
TOCOPHEROL (UNII: R0ZB2556P8)	
SOPHORA FLAVESCENS ROOT (UNII: IYR6K8KQ5K)	
HEXASODIUM PHYTATE (UNII: ZBX50UG81V)	
PALMITIC ACID (UNII: 2V16EO95H1)	
MENYANTHES TRIFOLIATA (UNII: 7H0QTZ446K)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70532-005-01	1 in 1 CARTON	04/03/2016	
1		50 mL 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	04/01/2016	

Labeler - IXXI S.A.S. (263290505)

Revised: 4/2016 IXXI S.A.S.