

PURE VELOCITY SUNSCREEN SPF 50- octinoxate cream
Guangzhou Haishi Biological Technology Co., Ltd.

Active Ingredients

OCTOCRYLENE
HOMOSALATE
OCTINOXATE

Purpose

SUNSCREEN

Uses

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

Stop Use

Rinse with wate to remove.
Stop use and ask doctor if rash occurs.

Do Not Use

Do not use on damaged or broken skin.

When Using

When using this product keep out of eyes.

Keep Out Of Reach Of Children

Keep Out Of Reach Of Children

Dosage & administration

Squeeze out an appropriate amount of Sunscreen and spread evenly on skin.

Inactive ingredients

Water
Glycerin
C12-15 Alkyl Benzoate
Dimethicone
Isododecane
Butyloctyl Salicylate
Propanediol
Stearic Acid
PEG-100 Stearate
Sorbitan Stearate
Niacinamide
Ceramide NP
Ceramide AP
Ceramide EOP
Sorbitan Isostearate
Carbomer
Cetearyl Alcohol
Ceteareth-20
Triethoxycaprylylsilane
Dimethiconol
Sodium Citrate
Sodium Lauroyl Lactylate
Sodium Dodecylbenzenesulfonate
Myristic Acid
Sodium Hyaluronate
Cholesterol
Aluminum Hydroxide
Palmitic Acid
Phenoxyethanol
Chlorphenesin
Hydroxyethyl Acrylate / Sodium Acryloyldimethyl Taurate Copolymer
Caprylyl Glycol
Citric Acid
Panthenol
Xanthan Gum
Phytosphingosine
Polyhydroxystearic Acid
Polysorbate 60
Ethylhexylglycerin



PURE VELOCITY SUNSCREEN SPF 50

octinoxate cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60771-0038
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	3.75 g in 100 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	6 g in 100 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	3 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60771-0038-1	75 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/30/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	09/30/2025	

Labeler - Guangzhou Haishi Biological Technology Co., Ltd. (421262738)

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Haishi Biological Technology Co., Ltd.		421262738	manufacture(60771-0038)

Revised: 9/2025

Guangzhou Haishi Biological Technology Co., Ltd.