

PURE VELOCITY SUNSCREEN SPF 50- ethylhexyl methoxycinnamate,zinc oxide liquid

Guangzhou Haishi Biological Technology Co., Ltd.

Active Ingredients

ETHYLHEXYL METHOXYCINNAMATE

ZINC OXIDE

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

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Dosage & administration

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

Inactive ingredients

WATER

PETROLATUM

ISOPROPYL PALMITATE

CETEARYL GLUCOSIDE

POLYDIMETHYLSILOXANE

HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER

POLYISOBUTYLENE

PEG-7 TRIMETHYLOLPROPANE COCONUT ETHER

SODIUM HYALURONATE

TOCOPHERYL ACETATE

POLYETHER-1

CITRIC ACID

OLEANOL POLYETHER-3 PHOSPHATE

PHENOXYETHANOL

BUTANEDIOL

IODOPROPYNYL BUTYLCARBAMATE

TRIETHOXYOCTYLSILANE



彩盒尺寸: 36x36x115mm

PURE VELOCITY SUNSCREEN SPF 50

ethylhexyl methoxycinnamate,zinc oxide liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60771-0031
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ETHYLHEXYL METHOXYCINNAMATE (UNII: 4Y5P7MUD51) (ETHYLHEXYL METHOXYCINNAMATE - UNII:4Y5P7MUD51)	ETHYLHEXYL METHOXYCINNAMATE	1.5 g in 100 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	3 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
PEG-7 TRIMETHYLOLPROPANE COCONUT ETHER (UNII: MVJ3AD73GG)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
POLYETHER-1 (2800 MPA.S) (UNII: 6QR7NJP3G8)	
.ALPHA.-BUTYLDIMETHYLSILYL-AND .OMEGA.-3-METHACRYLOXYPROPYLDIMETHYLSILYL-TERMINATED POLYDIMETHYLSILOXANE (10000 MW) (UNII: P7CQ6Y7QTS)	
CITRIC ACID (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0KO0R)	
POLYISOBUTYLENE (1000 MW) (UNII: 5XB3A63Y52)	
SODIUM HYALURONATE (UNII: YSE9PPT4TH)	
1,2-BUTANEDIOL (UNII: RUN0H01QEU)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60771-0031-1	30 g 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(60771-0031)

Revised: 9/2025

Guangzhou Haishi Biological Technology Co., Ltd.