

PURE VELOCITY TALLOW ZINC SUN BALM SPF 50- ethylhexyl salicylate,zinc oxide cream
Guangzhou Haishi Biological Technology Co., Ltd.

Active Ingredients

ETHYLHEXYL SALICYLATE
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

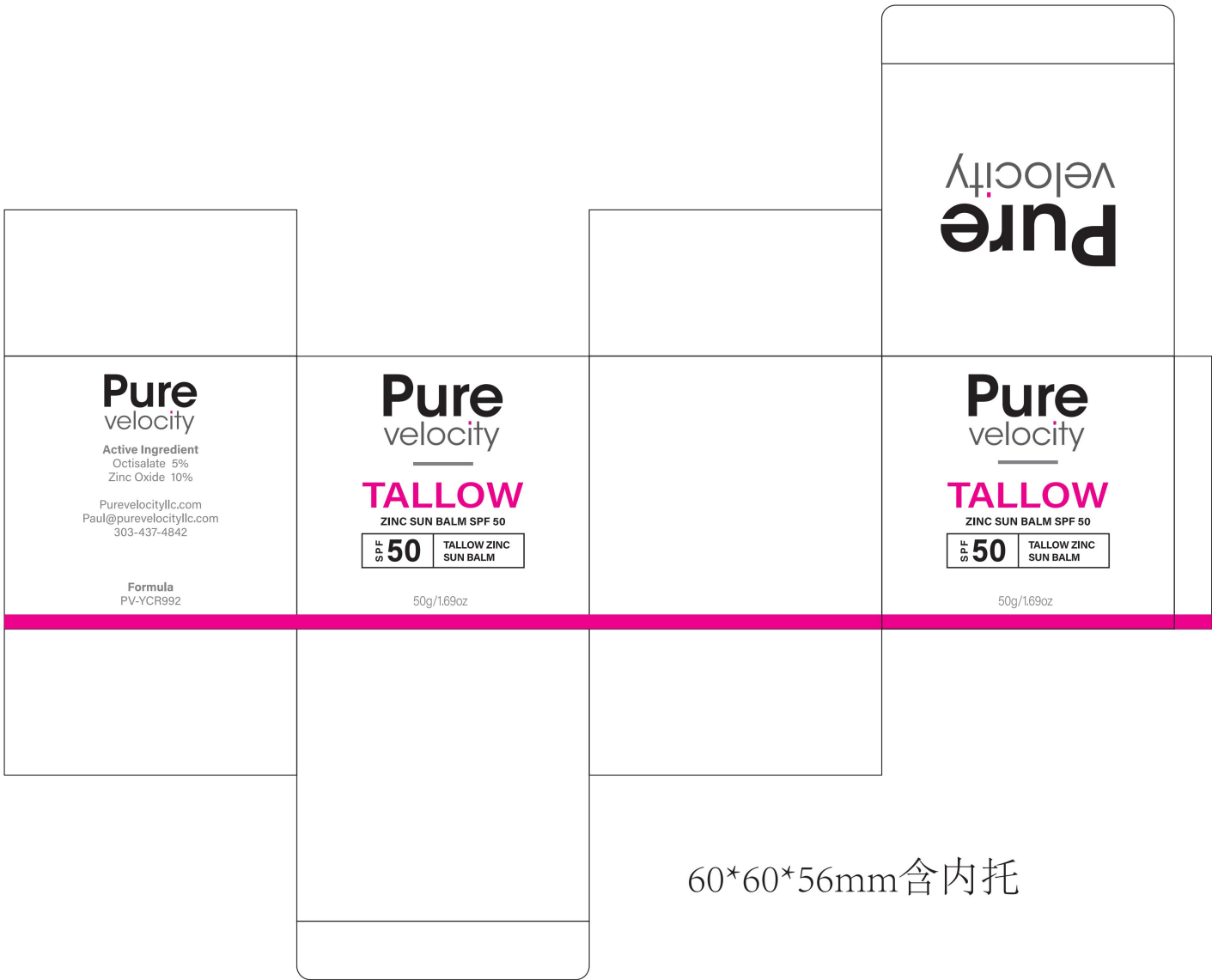
Keep Out Of Reach Of Children

Dosage & administration

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

Inactive ingredients

ETHYLHEXYL PALMITATE
ADEPS BOVIS
SIMMONDSIA CHINENSIS (JOJOBA) SEED OIL
OZOKERITE
CERA ALBA
DIMETHICONE
SILICA DIMETHYL SILYLATE
TOCOPHEROL
ROSA CANINA FRUIT OIL
PARFUM
BISABOLOL
WATER



60*60*56mm含内托

PURE VELOCITY TALLOW ZINC SUN BALM SPF 50
ethylhexyl salicylate,zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60771-0034
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	5 g in 100 g
ETHYLHEXYL SALICYLATE (UNII: 4X49Y0596W) (ETHYLHEXYL SALICYLATE - UNII:4X49Y0596W)	ETHYLHEXYL SALICYLATE	2.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PARFUMIDINE (UNII: A2Q2LHC6MA)	
WATER (UNII: 059QF0KO0R)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
SIMMONDSIA CHINENSIS (JOJOBA) SEED OIL (UNII: 724GKU717M)	
ABIES ALBA WHOLE (UNII: 62INA9V33X)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SILICA DIMETHYL SILYLATE (UNII: EU2PSP0G0W)	
ROSA CANINA FRUIT OIL (UNII: CR7307M3QZ)	
BISABOLOL (UNII: 24WE03BX2T)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
ADEPS BOVIS (UNII: 98HPY76U4W)	
BEESWAX (UNII: 2ZA36H0S2V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60771-0034-1	50 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(60771-0034)