CONJU PRINCESS UV SUN BLOCK - ethylhexyl methoxycinnamate cream
Conju Inc

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Drug Facts

active ingredients: Ethylhexyl methoxycinnamate, 4-methylbenzylidene camphor, butyl methoxydibenzoylmethane, titanium dioxide, zinc oxide

inactive ingredient: water, cyclopentasiloxane, dipropylene glycol, glycerin, dimethicone, hexyl laurate, sorbitan olivate, sorbitan sesquioleate, magnesium sulfate, stearic acid, hectorite, aluminum hydroxide, phenoxyethanol, propylparaben, methylparaben, disodium EDTA, fragrance

Recommended for sensitive skin that needs virtually total sunblock protection

keep out of reach of the children

Apply daily, with or without make-up to exposed areas

Avoid eye area. If contact occurs, rinse eyes thoroughly with water.
If following abnormal symptoms (red specks, swelling, itching) occurs after use, stop use and ask doctor.

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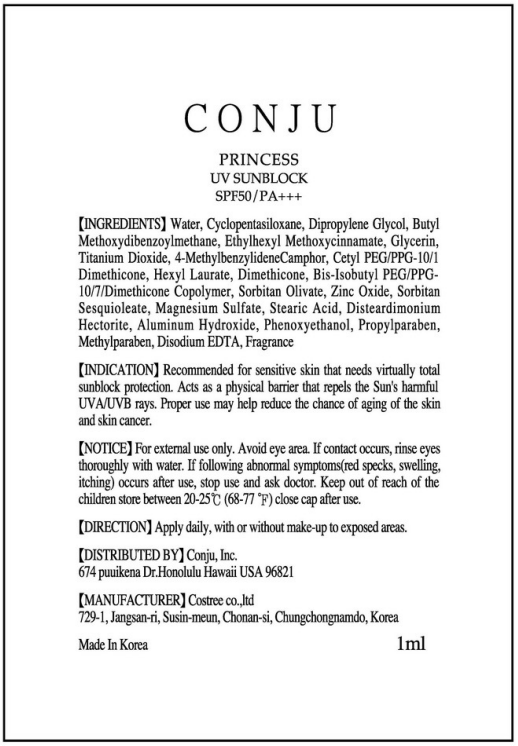

CONJU PRINCESS UV SUN BLOCK

ethylhexyl methoxycinnamate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59915-4001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	0.075 mL in 1 mL
ENZACAMENE (UNII: 8I3XWY40L9) (ENZACAMENE - UNII:8I3XWY40L9)	ENZACAMENE	0.05 mL in 1 mL
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.0249 mL in 1 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	0.03 mL in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.01 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
HEXYL LAURATE (UNII: 4CG9F9W01Q)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
HECTORITE (UNII: 08X4KI73EZ)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:59915-4001-1	1 mL in 1 POUCH		

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
unapproved drug other		12/18/2010	

Labeler - Conju Inc (012345486)

Registrant - Conju Inc (012345486)

Establishment

Name	Address	ID/FEI	Business Operations
Costree Co., Ltd		690345835	manufacture

Revised: 12/2010 Conju Inc