

**MISSHA TIME REVOLUTION WHITE CURE UV SUN PROTECTOR SPF50 PLUS PA PLUS PLUS PLUS- zinc oxide, octinoxate, octisalate cream
ABLE C&C CO., LTD.**

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.

ACTIVE INGREDIENT

ACTIVE INGREDIENTS: Zinc Oxide 10.24%, Ethylhexyl Methoxycinnamate 7.00%, Ethylhexyl Salicylate 2.00%

INACTIVE INGREDIENT

INACTIVE INGREDIENTS : Water, Cyclomethicone, Glycereth-26, Jasminum Officinale (Jasmine) Flower Extract, Chamomilla Recutita (Matricaria) Flower Extract, Freesia Alba Flower Extract, Niacinamide, Saussurea Involucrata Extract, Bellis Perennis (Daisy) Flower Extract, Lilium Candidum Flower Extract, Houttuynia Cordata Extract, Lauryl PEG-9 Polydimethylsiloxyethyl Dimethicone, Betaine, Gnaphalium Leontopodium Flower Extract, Sea Water, Echium Plantagineum Seed Oil, Cardiospermum Halicacabum Flower/Leaf/Vine Extract, Helianthus Annuus (Sunflower) Seed Oil Unsaponifiables, Cassia Alata Leaf Extract, Sigesbeckia Orientalis Extract, Radosia Rubescens Extract, Glucose, Methoxy PEG-45 Thiocetate, Buddleja Axillaris Leaf Extract, Fullerenes, Glyceryl Diisostearate, PEG-7 Dimethicone, C12-15 Alkyl Benzoate, Ethyl Hexanediol, Distearyltrimonium Chloride, PEG-30 Dipolyhydroxystearate, Polymethyl Methacrylate, Dimethicone/Methicone Copolymer, Beeswax, Sorbitan Oleate, Glyceryl Caprylate, Iron Oxides, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Xanthan Gum, Octyldodecanol, BHT, Methicone, Sodium C14-17 Alkyl Sec Sulfonate, Copper Sulfate, Polyacrylic Acid, Butylene Glycol, PVP, Fragrance, Butylphenyl Methylpropional, Eugenol

PURPOSE

PURPOSE: This combination of Cosmetic / OTC Drug can be used as a sun protection cream

WARNINGS

WARNING:

For external use only.

Do not swallow.

Keep out of reach of children.

Avoid contact with eyes.

Do not use on wounds or if you are allergic to any ingredients of this product.

KEEP OUT OF REACH OF CHILDREN

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INDICATIONS AND USAGE

INDICATION AND USAGE:

At the last step of basic skin care, apply an appropriate amount evenly over the face.

Pat lightly until fully absorbed.

Reapply during long exposure.

DOSAGE AND ADMINISTRATION

DOSAGE AND ADMINISTRATION: Take appropriate amount according to your skin condition and apply every morning and night.

PACKAGE LABEL. PRINCIPAL DISPLAY PANEL



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zinc oxide, octinoxate, octisalate cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:13733-019
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	3.89 g in 38 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	2.66 g in 38 g
OCTISALATE (UNII: 4X49 Y0596 W) (OCTISALATE - UNII:4X49 Y0596 W)	OCTISALATE	0.76 g in 38 g

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0K00R)	
Cyclomethicone (UNII: NMQ347994Z)	
Glycereth-26 (UNII: NNE56F2N14)	
Niacinamide (UNII: 25X51I8RD4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13733-019-01	38 g in 1 CARTON		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	03/01/2012	

Labeler - ABLE C&C CO., LTD. (689540284)

Registrant - ABLE C&C CO., LTD. (689540284)

Establishment

Name	Address	ID/FEI	Business Operations
ABLE C&C CO., LTD.		689540284	manufacture(13733-019)