LASER BLOCK 100 SUNBLOCK- octinoxate cream
Universal Cosmetic Co., Ltd

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, Zinc Oxide, 4-Methylbenzylidene camphor, Titanium Dioxide, Isoamyl p-Methoxycinnamate, Butyl Methoxydibenzoylmethane

Uses
■ Perfectly blocks UV rays without stimuli
■ Effective against
l PIH(Post Inflammatory Pigmentation)
l Photosensitive skins
l DNA destruction
■ Suitable for skin that underwent medical treatment
■ Helps keep moisturize

Warning
For external use only
When using this product
■ Avoid eye area. If contact occurs, rinse eyes thoroughly
■ If following abnormal symptoms occurs after use, stop use and ask doctor
l red specks, swelling, itching
Keep out of reach of the children

Direction
■ Before finishing skincare, rub a proper quantity over the whole face that underwent IPL or Laser therapy.

Other Information
■ store between 20-25 °C (68-77 °F)
■ avoid freezing and excessive heat above 40 °C (104 °F)
■ close cap after use.

LASER BLOCK 100 SUNBLOCK

octinoxate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52554-3001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	0.07 mL in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.05 mL in 1 mL
ENZACAMENE (UNII: 8I3XWY40L9) (ENZACAMENE - UNII:8I3XWY40L9)	ENZACAMENE	0.03 mL in 1 mL
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.02 mL in 1 mL
AMILOXATE (UNII: 376KTP06K8) (AMILOXATE - UNII:376KTP06K8)	AMILOXATE	0.02 mL in 1 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	0.005 mL in 1 mL

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52554-3001-1	50 mL in 1 TUBE		

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
unapproved drug other		08/04/2010	

Labeler - Universal Cosmetic Co., Ltd (557795012)

Revised: 8/2010 Universal Cosmetic Co., Ltd