TERADIA THE LIQUIDDIAMOND SUNSCREEN SPF50- ethylhexyl methoxycinnamate, homosalate, ethylhexyl salicylate cream
NanoDia Lab 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.

Drug Facts

Octinoxate, Homosalate, Octisalate

Niacinamide, Adenosine

Sunscreen, Anti-Wrinkle, Whitening

keep out of reach of the children

At the last step of skin care, take an appropriate amount and apply evenly on the face, neck and other areas exposed to UV radiation

1. Do not use in the following cases(Eczema and scalp wounds)
2.Side Effects
1)Due to the use of this druf if rash, irritation, itching and symptopms of hypersnesitivity occur dicontinue use and consult your phamacisr or doctor
3.General Precautions
1)If in contact with the eyes, wash out thoroughty with water If the symptoms are servere, seek medical advice immediately
2)This product is for exeternal use only. Do not use for internal use
4.Storage and handling precautions
1)If possible, avoid direct sunlight and store in cool and area of low humidity
2)In order to maintain the quality of the product and avoid misuse
3)Avoid placing the product near fire and store out in reach of children

for external use only

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ethylhexyl methoxycinnamate, homosalate, ethylhexyl salicylate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71483-0001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7 g in 100 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	6 g in 100 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	4.5 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ADENOSINE (UNII: K72T3FS567)	
NIACINAMIDE (UNII: 25X51I8RD4)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71483-0001-1	40 mL in 1 TUBE; Type 0: Not a Combination Product	12/20/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	12/20/2017	

Labeler - NanoDia Lab Co Ltd (693900835)

Registrant - NanoDia Lab Co Ltd (693900835)

Establishment

Name	Address	ID/FEI	Business Operations
NanoDia Lab Co Ltd		693900835	manufacture(71483-0001)

Revised: 12/2018 NanoDia Lab Co Ltd