

SNOW WHITE- niacinamide cream

Zenpia

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

niacinamide

Water, Cyclopentasiloxan, Cyclohexasiloxane, Glycerin, Mineral Oil, Propylene Glycol, Cetearyl Alcohol, Isododecane, Distearidimonium Hectorit, Propylene Carbonate, Titanium Dioxide, Mica, Polymethyl Methacrylate, Triethoxycaprylylsilane, Glyceryl Stearate, PEG-100 Stearate, Beeswax, Dimethicone, Stearic Acid, Polysorbate 60, Phenoxyethanol, Triethanolamine, Carbomer, Fragrance, Disodium EDTA

skin whitening

keep out or reach of the children

Take an appropriate amount with a spatula, spread on to your skin and tap it.

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 phamacistr 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

스노우 화이트 크림 단상자_국내용	
수량	
재질	CCP300
인쇄사양	2도
	 Pantone Cool gray 2c
	 Pantone Cool gray 10c



SNOW WHITE

niacinamide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70825-0001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X51I8 RD4) (NIACINAMIDE - UNII:25X51I8 RD4)	NIACINAMIDE	2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength

WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70825-0001-1	50 g in 1 JAR; Type 0: Not a Combination Product	07/05/2016	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		07/05/2015		

Labeler - Zenpia (557799448)

Registrant - Zenpia (557799448)

Establishment			
Name	Address	ID/FEI	Business Operations
Zenpia		557799448	label(70825-0001)

Establishment			
Name	Address	ID/FEI	Business Operations
EZEKIELCOSMETIC CO.,LTD		689851966	manufacture(70825-0001)