SNOW WHITE MILKY PACK- 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,Cyclohexasiloxan,Dimethicon,Cyclomethicon,Butylene Glycol,Cetyl PEG/PPG-10/1 Dimethicone,Dimethicone/Vinyl Dimethicone Crosspolymer,IsoDodecan,Disteardimonium Hectorite,Propylene Carbonate,Titanium Dioxide,Mica,Polymethyl Methacrylate,CI 77492,CI 77499,Triethoxycaprylylsilane,Sodium Hyaluronate,Sorbitan Sesquioleate,Ceresin,Magnesium Sulfate,Phenoxyethanol,Disodium EDTA,Fragrance

skin whitening

keep out or reach of the children

Apply onto cleaned face or body and gently massage it.

Wash off it with warm water after 2-3 minutes.

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

별색2도 / 무광라미

스노우 화이트 밀키 팩_리뉴얼	
수 량	
재 질	은페트지
인쇄사양	별색3도
	Pantone Cool gray 2c Pantone Cool gray 10c White

SNOW WHITE MILKY PACK

niacinamide cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70825-0008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	2 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70825-0008-1	200 g in 1 TUBE; Type 0: Not a Combination Product	07/11/2016	

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(70825-0008)

Establishment

Name	Address	ID/FEI	Business Operations
EZEKIELCOSMETIC CO.,LTD		689851966	manufacture(70825-0008)

Revised: 12/2018 Zenpia