PROTECTING NATURAL SHINE TONE UP SUN CUSHION REFILL- zinc oxide, titanium dioxide powder
In This Morning

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, adenosine, titanium dioxide, zinc oxide

water, butylene glycol, etc.

against UV, skin-brightening, alleviate wrinkles

keep out or reach of the children

apply moderate amount of this product evenly on skin

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

PROTECTING NATURAL SHINE TONE UP SUN CUSHION REFILL

zinc oxide, titanium dioxide powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71217-0045

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ADENOSINE (UNII: K72T3FS567) (ADENOSINE - UNII:K72T3FS567)	ADENOSINE	0.04 g in 100 g
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	2 g in 100 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	9.699515 g in 100 g
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	7.383 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:71217-0045-1	25 g in 1 PACKAGE; Type 0: Not a Combination Product	01/03/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/26/2019	

Labeler - In This Morning (694519423)

Registrant - In This Morning (694519423)

Establishment

Name	Address	ID/FEI	Business Operations
In This Morning		694519423	pack(71217-0045) , manufacture(71217-0045) , label(71217-0045)

Revised: 2/2020 In This Morning