KIDZGOO NATURAL SUNSCREEN- zinc oxide lotion
RENU LABORATORIES, INC.

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.

KidZGoo Natural Sunscreen

Active Ingredient

Zinc Oxide

Purpose

Sunscreen

Warnings

- **For external use only**
- **Do not use** on damaged or broken skin
- **When using this product,** keep out of eyes. Rinse with water to remove.
- **Stop use and ask a doctor** if rash occurs
- **Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

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855-KIDZGOO

Melearning Media LLC

Upper Dublin, PA

Directions

- apply liberally 15 minutes before sun exposure
- reapply;
  - after 80 minutes of swimming or sweating
  - immediately after towel drying
  - at least every 2 hours

Sun Protection Measures.

Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a broad spectrum value of SPF 15 or higher and other sun protection measures including

- limit time in the sun, especially from 10 a.m. - 2 p.m.
- wear long-sleeved shirts, pants, hats, and sunglasses
- children under 6 months of age: Ask a doctor

Uses

- helps prevent sunburn if used as directed with other sun protection measures
- (see **Directions**)
- Decreases the risk of skin cancer and early skin aging caused by the sun.

Inactive Ingredients:

Benzyl Alcohol, Caprylic/Capric Triglyceride, Cetearyl Alcohol, Cetearyl Olivate, Coco-Caprylate, Dehydroacetic Acid, Deionized Water, Isostearic Acid, Lecithin, Microcrystalline Cellulose, Panthenol, Polyglyceryl-3, Polyhydroxystearic Acid, Silica, Sodium Phytate, Sorbitan Olivate, Stearyl Octyldodecyl Citrate Crosspolymer, Tocopherol, Xantan Gum

KIDZGOO NATURAL SUNSCREEN

zinc oxide lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76348-650
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	98.56 g in 448 g

Inactive Ingredients	
Ingredient Name	**Strength**
STYRENE/ACRYLAMIDE COPOLYMER (MW 500000) (UNII: 5Z4DPO246A)	
PANTHENOL (UNII: WV9CM0O67Z)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
PHYTATE SODIUM (UNII: 88496G1ERL)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
STEARYL/OCTYLDODECYL CITRATE CROSSPOLYMER (UNII: PN88NW0KPK)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
COCO-CAPRYLATE (UNII: 4828G836N6)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:76348-650-16	448 g in 1 POUCH; Type 0: Not a Combination Product	07/30/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	07/30/2021	

Labeler - RENU LABORATORIES, INC. (945739449)

Registrant - RENU LABORATORIES, INC. (945739449)

Establishment

Name	Address	ID/FEI	Business Operations
RENU LABORATORIES, INC.		945739449	manufacture(76348-650)