QUARXERY QUARXERY PHYSICAL SUNSCREEN- quarxery physical sunscreen cream Guangzhou Meixi Biotechnology Co., Ltd.

84167-001

Active Ingredient

Zinc Oxide 16%

Purpose

Helps prevent sunburn

Use

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.

Warnings

For external use only

Do not use

Do not use on damaged or broken skin.

When Using

When using this product keep out of eyes. Rinse with wate to remove Stop use and ask doctor if rash occurs.

Stop Use

Stop use and ask doctor if rash occurs

Ask Doctor

Children under 6 months of age:

Keep Oot Of Reach Of Children

Keep out of reach of children. Ifswallowed, get medical help orcontact Poison Control Center right away.

Directions

Apply liberally and evenly 15 minutes prior to sun exposure. Reapply after 40 minutes of swimming or sweating immediately aftertowel drying "at least every 2 hours.

Other information

Protect this product from excessive heat and direct sun.

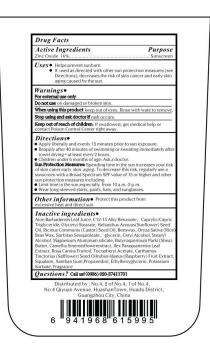
Inactive ingredients

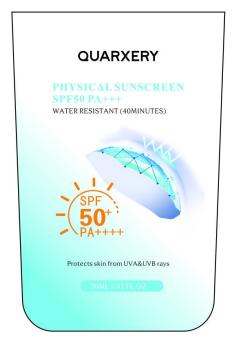
Aloe Barbadensis Leaf Juice, C12-15 Alky Benzoate Caprylic/Capric Triglyceride, Glyceryi Stearate, Helianthus Annuus (Sunflower) Seed Oil, Ricinus Communis (Castor) Seed Oil, Beeswax, Oryza Sativa (Rice) Bran Wax, Sorbitan Sesquioleate glycerin, Cetyi Alcohol, Stearyl Alcohol. Magnesium Aluminum silicate, Butyrospermum Parki (Shea) Butter, Camellia Sinensis flower extract, Ilex Paraguariensis Leaf Extract, Rosa Canina Fruitoil, Tocopheryi Acetate, Carthamus Tinctorius (Safflower) Seed, Oilrubus Idaeus (Raspberry) Fruit Extract, Squalane, Xanthan Gum, Propanediol, Ethylhexy glycerin, Potassium Sorbate, Fragrance

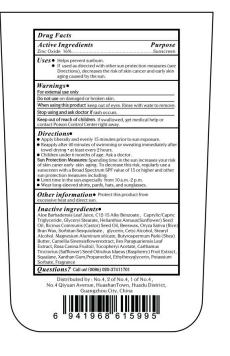
Questions

Call us!(0086)020-37411701

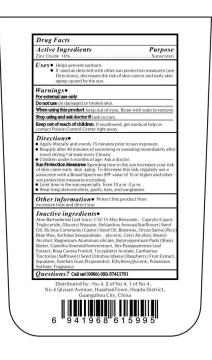
PRINCIPAL DISPLAY PANEL



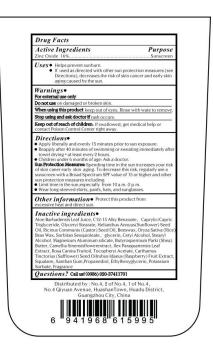




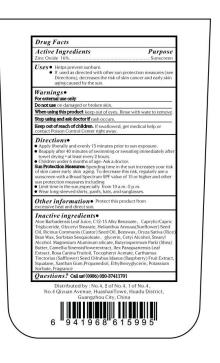




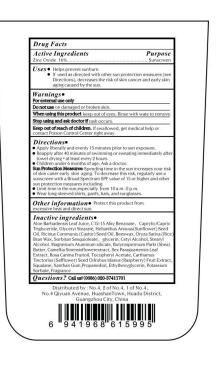




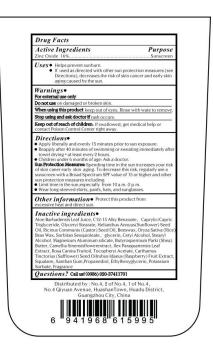


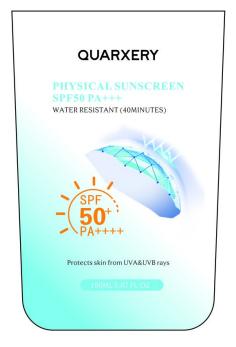


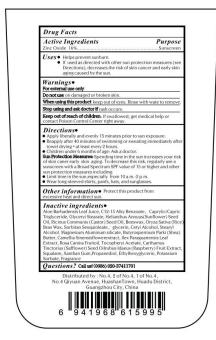














QUARXERY QUARXERY PHYSICAL SUNSCREEN

quarxery physical sunscreen cream

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:84167-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)

ZINC OXIDE

16 g in 100 mL

Inactive Ingredients Ingredient Name Strength RICE BRAN (UNII: R60QEP13IC) CAMELLIA SINENSIS FLOWER (UNII: 9I2BJY2J17) SAFFLOWER SEED (UNII: 8FCL6A31TX)

XANTHAN GUM (UNII: TTV12P4NEE)
CASTOR OIL (UNII: D5340Y2I9G)

GLYCERIN (UNII: PDC6A3C0OX)	
PROPANEDIOL (UNII: 5965N8W85T)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
RASPBERRY (UNII: 4N14V5R27W)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALLYL BENZOATE (UNII: AQY5543KFE)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
SQUALANE (UNII: GW89575KF9)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)	
ROSA CANINA FRUIT OIL (UNII: CR7307M3QZ)	
SHEA BUTTER (UNII: K49155WL9Y)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
FRAGRANCE 13576 (UNII: 5EM498GW35)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:84167-001- 01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024			
2	NDC:84167-001- 02	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024			
3	NDC:84167-001- 03	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024			
4	NDC:84167-001- 04	90 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024			
5	NDC:84167-001- 05	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024			
6	NDC:84167-001- 06	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024			
7	NDC:84167-001- 07	150 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024			
8	NDC:84167-001- 08	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	M020	03/13/2024				

Establishment						
Name	Address	ID/FEI	Business Operations			
Guangzhou Meixi Biotechnology Co., Ltd.		625724571	label(84167-001), manufacture(84167-001)			

Revised: 3/2024 Guangzhou Meixi Biotechnology Co., Ltd.