BRIGHTER DAY BROAD SPECTRUM SPF 30 SUNSCREEN- zinc oxide lotion
SQN Lab

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.

SQN - BRIGHTER DAY BROAD SPECTRUM SPF30 (82495-101)

ACTIVE INGREDIENT

ZINC OXIDE

PURPOSE

SUNSCREEN

USE

HELPS PREVENT SUNBURN

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 PRODUCT IS SWALLOWED, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER RIGHT AWAY.

DIRECTIONS

- APPLY LIBERALLY 15 MINUTES BEFORE SUN EXPOSURE.
- CHILDREN UNDER 6 MONTHS OF AGE: ASK A DOCTOR.
- USE A WATER-RESISTANT SUNSCREEN IF SWIMMING OR SWEATING.
- REAPPLY AT LEAST EVERY 2 HOURS.

- SUN PROTECTION MEASURES.
  SPENDING TIME IN THE SUN INCREASES YOUR RISK OF SKIN CANCER AND EARLY AGING. TO DECREASE THE RISK, REGULARLY USE A SUNSCREEN WITH A BROAD SPECTRUM SPF VALUE OF 15 OR HIGHER AND OTHER SUN PROTECTION MEASURES INCLUDING:
- LIMIT TIME IN THE SUN, ESPECIALLY FROM 10:00 A.M. - 2 P.M.
- WEAR LONG-SLEEVED SHIRTS, PANTS, HATS, AND SUNGLASSES.

INACTIVE INGREDIENTS

WATER, BUTYLENE GLYCOL, CAPRYLIC/CAPRIC TRIGLYCERIDE, DIETHYLHEXYL 2,6-NAPHTHALATE, HYDROXYETHYL ACRYLATE/SODIUM, ACRYLOYLDIMETHYL TAURATE COPOLYMER, SODIUM OLIVATE, XYLITYLGLUCOSIDE, ANHYDROXYLITOL, TRIETHOXYCAPRYLYLSILANE, XYLITOL,PENTYLENE GLYCOL, CITRIC ACID, TOCOPHEROL, CAMELLIA SINENSIS LEAF OIL, DIMETHICONE, PHENOXYETHANOL, ETHYLHEXYLGLYCERIN, IRON OXIDES.

OTHER INFORMATION

- PROTECT THE PRODUCT IN THIS CONTAINER FROM EXCESSIVE HEAT AND DIRECT SUN.

BRIGHTER DAY BROAD SPECTRUM SPF 30 SUNSCREEN

zinc oxide lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82495-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	10 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
DIETHYLHEXYL 2,6-NAPHTHALATE (UNII: I0DQJ7YGXM)	
TOCOPHEROL (UNII: R0ZB2556P8)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
WATER (UNII: 059QF0KO0R)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
SODIUM OLEATE (UNII: 399SL044HN)	
XYLITYLGLUCOSIDE (UNII: O0IEZ166FB)	
ANHYDROXYLITOL (UNII: 8XWR7NN42F)	
XYLITOL (UNII: VCQ006KQ1E)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
TEA LEAF OIL (UNII: VC855RRT77)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82495-101-11	1 in 1 BOX	12/20/2021	
1		50 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	12/20/2021	

Labeler - SQN Lab (118483828)

Revised: 1/2022 SQN Lab