

BABIES KIDSSUN PROTECTION- mineral sunscreen powder
PANERGYTEK USA INC

Titanium dioxide-19.5%

Zinc oxide-24%

Sunscreen

Helps prevent sunburn.If used as directed with other sunprotection measures(see Directions)decreases the risk of skin cancer andearly skin aging caused by the sun.

For external use only.Do not use on damaged or broken skinWhen using this product keep out ofeyes. Rinse with water to remove.Stop use and ask a doctor if rash occursKeep out of reach of children. Ifproducts swallowed. get medical helpor contact a Poison Control Center rightaway.

Apply liberally 15 minutes before sunexposure.

Reapply: After 80 minutes ofswimming or sweating. immediatelyafter towel drying.At least every 2 hours.Sun Protection Measures.Spending time in the sun increasesyour risk of skin cancer and early skinaging. To decrease this risk, regularlyuse a sunscreen with a broadspectrum SPF value of 15 or higher.And other sun protection measuresincluding: Limit time in the sun,especially from 10am-2pm. wearlong-sleeved shirts, pants, hats andsunglasses. Children under 6months of age: ask a doctor

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Protect this product in this container from excessive heat and direct sun.

Vinyl Dimethicone/MethiconeSilsesquioxane Crosspolymer.

Aluminum hydroxide.

LavandulaAngustifolia (Lavender) Oil,

Mica.

Lophatherum Gracile Leaf Power,

Dimethicone,

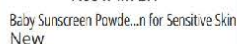
Triethoxycoprylsilane

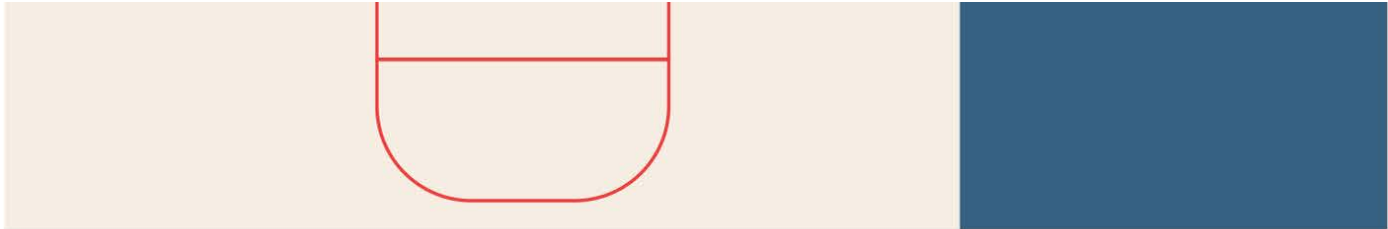
.Stearic Acid,

Tocopherol(Vitamin E),

C1 77492,

Sodium Benzoate
,Potassium Sorbate





BABIES KIDSSUN PROTECTION

mineral sunscreen powder

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85950-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	240 mg in 1 g
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	195 mg in 1 g

Inactive Ingredients	
Ingredient Name	Strength
SODIUM BENZOATE (UNII: OJ245FE5EU)	
VINYL DIMETHICONE/METHICONE SILSESQUIOXANE CROSSPOLYMER (UNII: 9NH1UDD2RR)	
POTASSIUM SORBATE (UNII: 1VPU26JZ Z4)	
MICA (UNII: V8A1AW0880)	
CI 77491 (UNII: 1K09F3G675)	
LAVANDULA ANGUSTIFOLIA (LAVENDER) OIL (UNII: ZBP1YXW0H8)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
STEARIC ACID (UNII: 4ELV7Z 65AP)	
TOCOPHEROL (UNII: R0ZB2556P8)	
LOPHATHERUM GRACILE LEAF (UNII: R417138GE9)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
CI 77492 (UNII: EX438O2MRT)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85950-001-01	5 g in 1 BOTTLE; Type 0: Not a Combination Product	07/20/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC Monograph Drug	M020	07/20/2025	
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Labeler - PANERGYTEK USA INC (122142088)

Registrant - PANERGYTEK USA INC (122142088)

Revised: 7/2025

PANERGYTEK USA INC