BABY PURE AND GENTLE SPF 60 SUNSCREEN- titanium dioxide - 6.00% zinc oxide - 4.70% stick CVS

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.

CVS Health

Drug Facts

Active ingredients	Purpose	
Titanium Dioxide - 6.00%	Sunscreen	
Zinc Oxide - 4.70%	Sunscreen	

Uses

helps prevent sunburn

Warnings For external use only

Do not use on damage 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 the children. If swallowed, get medical help or contact a Poison Control Center right away

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 SPF of 15 or higher and other sun protection measures including:

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

Inactive ingredients

Aluminum Hydroxide, Beeswax, BHT, Butyloctyl Salicylate, C12-15 Alkyl Benzoate, Cetyl Alcohol, Dimethicone, Euphorbia Cerifera (Candelilla) Wax, Isostearic Acid, Neopentyl Glycol Diethylhexanoate, Ozokerite, Paraffin, Polyethylene, Stearic Acid, Triethoxycaprylylsilane

Other Information

• protect this product from excessive heat and direct sun



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and direct sun. Inactive ingre Aluminum Hydroxide, Benzoate, Cetyl Alcoh Isostearic Acid, Neope Polyethylene, Stearic	edients Beeswax, BHT, Buty ol, Dimethicone, Eup entyl Glycol Diethylh Acid, Triethoxycapry	loctyl Salicylat horbia Cerifera exanoate, Ozol lylsilane.	e, C12-15 Alkyl a (Candelilla) Wax
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titanium dioxide - 6.00% zinc oxide - 4.70% stick

Product Information	n				
Product T ype	HUMAN OTC DRUG	Item Code ((Source)	NDC:69842-333	
Route of Administration	on TOPICAL				
Active Ingredient/A	Active Moiety				
	Ingredient Name		Basis of Stre	ength Strength	
Titanium Dioxide (UNII:	15FIX9V2JP) (TITANIUM DIOXIDE ·	· UNII:15FIX9V2JP)	Titanium Dioxide	e 6 g in 100 g	
Zinc Oxide (UNII: SOI2L	OH54Z) (ZINC CATION - UNII:13S1S	8SF37)	ZINC CATION	4.7 g in 100 g	
Inactive Ingredien	ts				
	Ingredient N	ame		Strength	
Aluminum Hydroxide (U	JNII: 5QB0T2IUN0)				
YELLOW WAX (UNII: 22	ZA36H0S2V)				
BUTYLATED HYDRO XY	YTOLUENE (UNII: 1P9 D0 Z171K)				
Butyloctyl Salicylate (U	NII: 2EH13UN8D3)				
ALKYL (C12-15) BENZ	DATE (UNII: A9EJ3J61HQ)				
Cetyl Alcohol (UNII: 936	JST6JCN)				
Dimethicone (UNII: 92R	J3N3Y1O)				
CANDELILLA WAX (UN	II: WL0328HX19)				
Isostearic Acid (UNII: X3	33R8U0062)				
Neopentyl Glycol Dieth	ylhexanoate (UNII: U68ZV6W62C)				
CERESIN (UNII: Q1LS2U	JO3A)				
Paraffin (UNII: 1900E3H	2ZE)				
HIGH DENSITY POLYE	THYLENE (UNII: UG00KM4WR7)				
Stearic Acid (UNII: 4ELV	77Z65AP)				
Triethoxycaprylylsilan	e (UNII: LDC331P08E)				
Packaging					
# Item Code	Package Descripti	on Mai	keting Start Date	Marketing End Dat	
1 NDC:69842-333-01 0	.47 g in 1 TUBE; Type 0: Not a Coml	Dination Product 02/13	3/2013		
Marketing Info	rmation				
	Application Number or Mon	agraph Citation M	arketing Start Date	Marketing End Da	
Marketing Category	Application Number of Mon		arketing Start Date	Marketing End Du	

Labeler - CVS (062312574)

Registrant - Product Quest Mfg (927768135)

Establishment

927768135

Revised: 4/2018