AFTER BITE KIDS- sodium bicarbonate cream Adventure Ready Brands

After Bite Kids

Drug Facts

Active Ingredient

Sodium Bicarbonate (Baking Soda) 5%

Purpose

Skin Protectant

Uses

Temporarily protects and helps relieve minor skin irritation and itching due to

- Insect bites
- Poison ivy, oak, or sumac

Warnings

for external use only..

When using this product

do not get into eyes

Stop use and ask a doctor if

- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few days

Keep Out of Reach of Children.

If swallowed, get medical help or contact a Poison Control Center right away. If in eyes flush with water for 15 minutes and call a doctor.

Directions

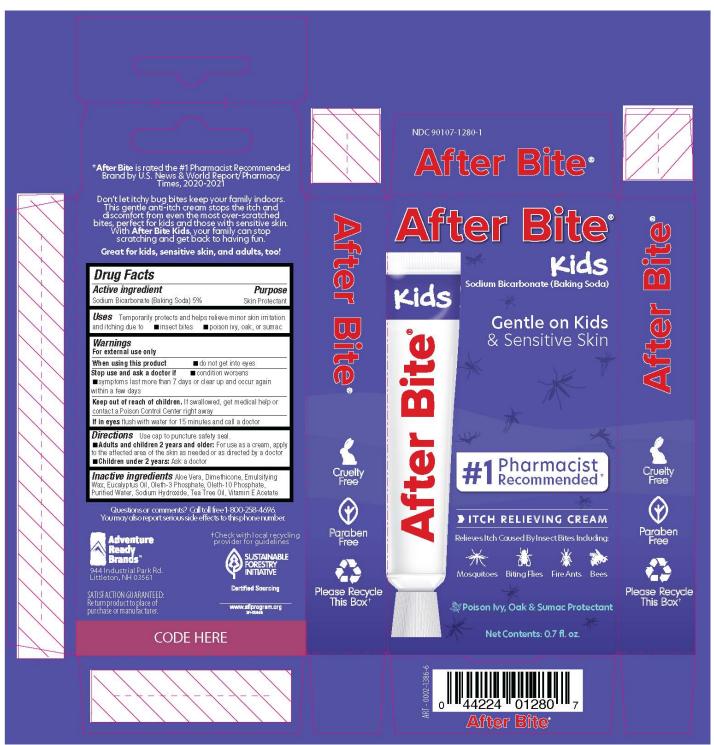
Use cap to puncture safety seal.

- Adults and children 2 years and older for use as a cream, apply to the affected area of the skin as needed or as directed by a doctor
- Children under 2 years ask a doctor

Inactive Ingredients

Aloe Vera, Dimethicone, Emulsifying Wax, Eucalyptus Oil, Oleth-3 Phosphate, Oleth-10 Phosphate, Purified Water, Sodium Hydroxide, Tea Tree Oil and Vitamin E Acetate

Package Label



AFTER BITE KIDS

sodium bicarbonate cream

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:90107-1280	
Route of Administration	TOPICAL			

	Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength	
l	SODIUM BICARBONATE (UNII: 8MDF5V39QO) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM BICARBONATE	1 g in 20 g	

Inactive Ingredients			
Ingredient Name	Strength		
ALOE VERA LEAF (UNII: ZY81Z83H0X)	1.1 g in 20 g		
DIMETHICONE (UNII: 92RU3N3Y1O)	0.1 g in 20 g		
EUCALYPTUS OIL (UNII: 2R04ONI662)	0.26 g in 20 g		
OLETH-3 PHOSPHATE (UNII: 8Q0Z18J1VL)	0.5 g in 20 g		
OLETH-10 PHOSPHATE (UNII: Q95361F4VS)	0.28 g in 20 g		
WATER (UNII: 059QF0KO0R)	14.1 g in 20 g		
SODIUM HYDROXIDE (UNII: 55X04QC32I)	0.24 g in 20 g		
TEA TREE OIL (UNII: VIF565UC2G)	0.2 g in 20 g		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	0.1 g in 20 g		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:90107- 1280-1	1 in 1 BOX	09/01/2020		
1		20 g 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 Drug	M016	09/01/2020		

Labeler - Adventure Ready Brands (064437304)

Revised: 2/2024 Adventure Ready Brands