KROGER BABY SUNSCREEN SPF 50- avobenzone, homosalate ,octisalate, octocrylene spray THE KROGER COMPANY

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.

Kroger Baby Sunscreen Spray SPF 50

Active Ingredients

Avobenzone 3.0% Homosalate 13.0% Octisalate 5.0% Octocrylene 2.0%

Purpose

Sunscreen

Uses

• helps prevent sunburn

• 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

• on damaged or broken skin.

When using this product

 keep out of eyes. Rinse with water to remove.
Keep away from face to avoid breathing it.
do not puncture or incinerate. Contents under pressure.
do not store at temperatures above 120° F.

Stop use and ask a doctor if

rash occurs.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

Directions

- Spray liberally and spread evenly by hand 15 minutes before sun exposure
- Hold can 4-6 inches away from the skin to apply
- Do not spray directly into face. Spray into hands then apply to face
- Do not apply in windy conditions
- Use in a well-ventilated area
- reapply:
- after 80 minutes of swimming or sweating
- immediately after towel drying
- at least every 2 hours
- children under 6 months of age: Ask a doctor

• **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 with a Broa Spectrum SPF value 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-sleeved shirts, pants, hats, and sunglasses

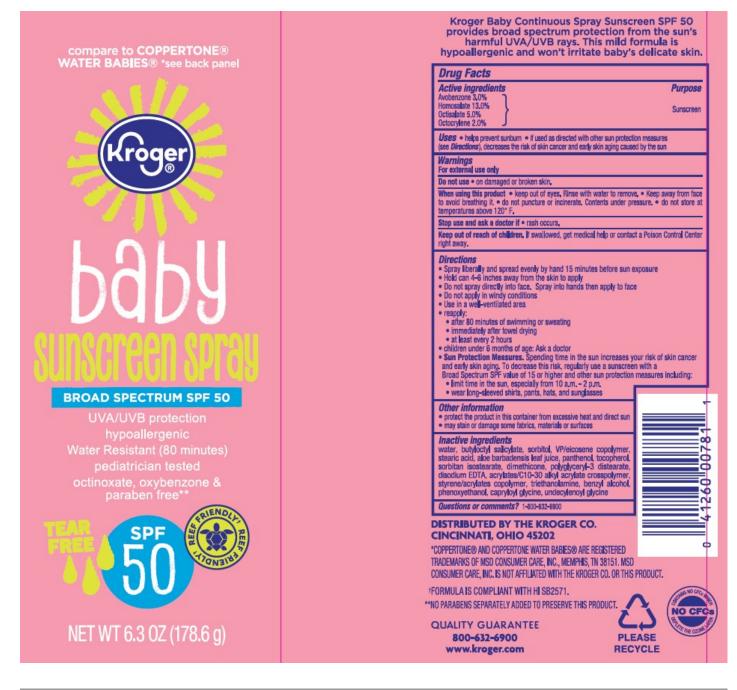
Other information

- protect the product in this container from excessive heat and direct sun
- may stain or damage some fabrics, materials or surfaces

Inactive Ingredients

water, butyloctyl salicylate, sorbitol, VP/eicosene copolymer, stearic acid, aloe barbadensis leaf juice, panthenol, tocopherol, sorbitan isostearate, dimethicone, polyglyceryl-3 distearate, disodium EDTA, acrylates/C10-30 alkyl acrylate crosspolymer, styrene/acrylates copolymer, triethanolamine, benzyl alcohol, phenoxyethanol, capryloyl glycine, undecylenoyl glycine

Label



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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:30142-237
Route of Administration	TOPICAL		

Active Ingredie	nt/Active Moiety			
	Ingredient Name	Basis of Str	ength Strength	
AVOBENZONE (UNII:	G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2	NOX) AVOBENZONE	30 mg in 1 g	
HOMOSALATE (UNII:	V06SV4M95S) (HOMOSALATE - UNII:V06SV4M9	5S) HOMOSALATE	HOMOSALATE 130 mg in 1	
OCTISALATE (UNII: 4	X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	4X49Y0596W) OCTISALATE 50		
OCTOCRYLENE (UNI	I: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WG	F6WM) OCTOCRYLENE	20 mg in 1 g	
Inactive Ingred	lients			
_	Ingredient Name		Strengt	
POLYGLYCERYL-3 D	DISTEARATE (UNII: ZI1LK470XV)		Jui Chige	
	(UNII: 7FLD91C86K)			
	YMER TYPE A (ALLYL PENTAERYTHRITOL CI	ROSSLINKED) (UNII: 71DD	5V995L)	
	IDE COPOLYMER (500000 MW) (UNII: 5Z4DF		,	
TROLAMINE (UNII: 90		,		
BENZYL ALCOHOL (
PHENOXYETHANOL	(UNII: HIE492ZZ3T)			
CAPRYLOYL GLYCIN	IE (UNII: 8TY5Y042NJ)			
UNDECYLENOYL GL	YCINE (UNII: 4D20464K2J)			
WATER (UNII: 059QF	0KO0R)			
BUTYLOCTYL SALIC	YLATE (UNII: 2EH13UN8D3)			
SORBITOL (UNII: 506	5T60A25R)			
VINYLPYRROLIDON	E/EICOSENE COPOLYMER (UNII: 035MV9S1C3)		
STEARIC ACID (UNII:	4ELV7Z65AP)			
ALOE VERA LEAF (U	NII: ZY81Z83H0X)			
PANTHENOL (UNII: W	W9CM0067Z)			
ALPHATOCOPHER	ROL, DL- (UNII: 7QWA1RIO01)			
SORBITAN ISOSTEA	RATE (UNII: 01S2G2C1E4)			
DIMETHICONE (UNII	: 92RU3N3Y1O)			
Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	178 g in 1 CAN; Type 0: Not a Combination Product	11/25/2019		
Marketing I	nformation			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	M020	11/25/2019		

Labeler - THE KROGER COMPANY (006999528)