

H.E.B BABY- spf 50 sunscreen aerosol, spray

H.E.B

H-E-B SOLUTIONS SUNSCREEN BABY SPF 50 LOTION

Active Ingredients

Avobenzone 3.0%, Homosalate 10.0%, Octisalate 4.5%, Octocrylene 8.0%

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.

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

- apply liberally 15 minutes before sun exposure
- 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 broad-spectrum 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

Other Information

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

☐ Inactive Ingredient(s)

water, butyloctyl salicylate, butylene glycol, glyceryl stearate, microcrystalline cellulose, behenyl alcohol, butylated PVP, stearic acid, lauryl alcohol, myristyl alcohol, cetyl alcohol, caprylic/capric triglyceride, tocopherol, sodium ascorbyl phosphate, cellulose gum, disodium EDTA, benzoic acid, benzoic alcohol, sorbic acid.

Questions or Comments?

Call toll free 1-800-527-7731

H-E-B SOLUTIONS SUNSCREEN BABY SPF 50 LOTION

8 FL OZ (237mL)

NDC 37808-913-12



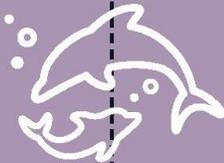
SOLUTIONS™ SUNSCREEN

BABY 50

Oxybenzone Free Sunscreen

UVA/UVB Protection
Water Resistant
(80 minutes)
Paraben Free

BROAD SPECTRUM SPF 50



8 FL OZ (237 mL)

Dermatologist Tested
Oil Free • PABA Free
Retinyl Palmitate Free • Fragrance Free
Hypoallergenic
Contains Vitamin E

Drug Facts

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Avobenzone 3% Homosalate 10% Octisalate 4.5% Octocrylene 8%	Sunscreen

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- at least every 2 hours

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• **Sun Protection Measures.** Spending time in the sun increases your risk of skin cancer and early aging. To decrease this risk, regularly use a sunscreen with a broad-spectrum SPF of 15 or higher and other sun protection measures including:

- limit time in the sun, especially from 10am-2pm
- wear long-sleeve shirts, pants, hats, and sunglasses.

Other Information • protect this product from excessive heat and direct sun
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Inactive Ingredient(s)

water, butyloctyl salicylate, butylene glycol, glyceryl stearate, microcrystalline cellulose, behenyl alcohol, butylated PVP, stearic acid, lauryl alcohol, myristyl alcohol, cetyl alcohol, caprylic/capric triglyceride, tocopherol, sodium ascorbyl phosphate, cellulose gum, disodium EDTA, benzoic acid, benzoic alcohol, sorbic acid.

Questions or Comments? Call toll free 1-800-527-7731

MADE WITH PRIDE AND CARE FOR H-E-B®,
SAN ANTONIO, TEXAS 78204



9569-1811



H.E.B BABY

spf 50 sunscreen aerosol, spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-913
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZ ONE	30 mg in 1 mL

HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	100 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	45 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	80 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DOCOSANOL (UNII: 9G1OE216XY)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
MYRISTYL ALCOHOL (UNII: V42034O9PU)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SORBIC ACID (UNII: X045WJ989B)	
LAURYL ALCOHOL (UNII: 178A96NLP2)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
SODIUM ASCORBYL PHOSPHATE (UNII: 836SJG51DR)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
TOCOPHEROL (UNII: R0ZB2556P8)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-913-12	237 mL in 1 TUBE; Type 0: Not a Combination Product	02/05/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	02/05/2019	

Labeler - H.E.B (007924756)