# HEB SHIMMER SUNSCREEN SPF 50- avobenzone, homosalate, octisalate, octocrylene lotion H.E.B

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.

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#### **HEB Shimmer Sunscreen SPF 50 Lotion**

### **∏Active Ingredients**

Acobenzone 3.0%, Homosalate 15.0%, Octisalate 5.0%, Octocrylene 7.0%

#### **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 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 aging. To decrease this risk, regularly use a sunscreen with a Broad Spectrum value of 15 or higher and other sun protection measures including:
- limit time in the sun, especially from 10am-2pm
- 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 Ingredient(s)

water, tridecyl salicylate, sorbitol, stearic acid, triethanolamine, aluminum starch octenylsuccinate, aloe barbadensis leaf juice, polyethyloxazoline, carbomer, dimethicone, tocopherol, polyglyceryl-3 distearate, caprylyl glycol, phenoxyethanol, ethylhexylglycerin, sorbitan isostearate, benzyl alcohol, fragrance, polyethylene terephthalate, polyurethane-33, aluminum powder, red 7, yellow 5, titanium dioxide, iron oxides, mica

#### Label



Dermatologist Tested · Hypoallergenic Oil Free · PABA Free · Retinyl Palmitate Free Contains Vitamin E

# Drug Facts

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Questions or comments? Call to free 1-800-527-7731

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





3089-2002



# **HEB SHIMMER SUNSCREEN SPF 50**

avobenzone, homosalate, octisalate, octocrylene lotion

#### **Product Information**

NDC:37808-041 **Product Type** HUMAN OTC DRUG Item Code (Source)

**Route of Administration TOPICAL** 

Active Ingredient/Active Moiety				
Ingredient Name	<b>Basis of Strength</b>	Strength		
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL		
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII: V06SV4M95S)	HOMOSALATE	150 mg in 1 mL		
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL		
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	70 mg in 1 mL		

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: 19PJ0O6294)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHATOCOPHEROL, DL- (UNII: 7QWA1RIO01)	
MICA (UNII: V8A1AW0880)	
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLOXAZOLINE (5000 MW) (UNII: HNX7574GTX)	
POLYGLYCERYL-3 DISTEARATE (UNII: ZI1LK470XV)	
TRIDECYL SALICYLATE (UNII: AZ Q08K38Z1)	
POLYETHYLENE TEREPHTHALATE (INTRINSIC VISCOSITY 1.00-2.00) (UNII: KJ6GWA9DJ0)	
ALUMINUM (UNII: CPD4NFA903)	
D&C RED NO. 7 (UNII: ECWOLZ 41X8)	
DIMETHICONE (UNII: 92RU3N3Y10)	
TROLAMINE (UNII: 903K93S3TK)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
SORBITOL (UNII: 506T60A25R)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:37808-041- 09	89 mL in 1 TUBE; Type 0: Not a Combination Product	05/08/2020		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	M020	05/08/2020		

# **Labeler -** H.E.B (007924756)

Revised: 2/2023 H.E.B