# HEB SPF 50 POMEGRANATE SUNSCREEN- 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 SPF 50 Pomegranate Sunscreen 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 protetion 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 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 Ingredient(s)**

water, polyester-7, neopentyl glycol diheptanoate, olive oil polyglyceryl-6 esters, glycerin, styrene/acrylates copolymer, aluminum starch octenylsuccinate, saccharomyces/podophyllum peltatum ferment filtrate, butylene glycol, silica, phenoxyethanol, isododecane, sodium stearoyl lactylate, cetearyl alcohol, cetearyl olivate, sorbitan olivate, cetyl palmitate, sorbitan palmitate, ethylhexylglycerin, glyceryl stearate, arachidyl alcohol, ethylhexylglycerin, acrylates/C10-30 alkyl acrylate crosspolymer, tocopherol (vitamin E), potassium hydroxide, sodium ascorbyl phosphate, fragrance.

#### Label



#### **HEB SPF 50 POMEGRANATE SUNSCREEN**

avobenzone, homosalate, octisalate, octocrylene lotion

# Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:37808-043 Route of Administration TOPICAL

<b>Active</b>	Ingredient/Active	Moiety
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Ingredient Name Basis of Strength Strength

AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	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)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: 19PJ0O6294)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PEG-100 STEARATE (UNII: YD01N1999R)	
.ALPHATOCOPHEROL, DL- (UNII: 7QWA1RIO01)	
POLYESTER-7 (UNII: 0841698D2F)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	
ISODODECANE (UNII: A8289P68Y2)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
CETYL PALMITATE (UNII: 5ZA2S6B08X)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
PODOPHYLLUM (UNII: 2S713A4VP3)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
OLIVE OIL POLYGLYCERYL-6 ESTERS (UNII: 4KDO9AFM9I)	
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
SODIUM STEAROYL LACTYLATE (UNII: IN99IT31LN)	
SORBITAN MONOPALMITATE (UNII: 77K6Z421KU)	
BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER (UNII: V5RS026Q0H)	

l	P	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:37808-043- 12	237 mL in 1 TUBE; Type 0: Not a Combination Product	03/10/2020		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M020	03/10/2020	

Revised: 2/2023 H.E.B