

**SEA STAR SPARKLE SPF 50 GLITTER SUNSCREEN COTTON CANDY-
homosalate, octocrylene, octisalate, zinc oxide cream
MJ Products Association, LLC**

SEA STAR SPARKLE SPF-50 Glitter Sunscreen Cotton Candy

Drug Facts

Active Ingredients

Homosalate 10.00%

Octocrylene 5.00%

Octisalate 5.00%

Zinc Oxide 4.00%

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 consult physician

if irritation or rash develops.

Keep out of reach of children.

If product is swallowed, get medical attention or contact poison control center immediately.

Directions

- Apply liberally 15 minutes before sun exposure.
- At least every two-hours.
- Immediately after swimming, sweating or towel drying. Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use sunscreen with broad spectrum protection and on SPF rating of 15 or higher, along with other sun protection measures including: Limit time in the sun, especially between 10am and 2pm. **SUN PROTECTION MEASURES:**
- Wear long sleeve shirts, pants, hats, and sunglasses.
- For children under 6 months, consult a physician.

REAPPLY:

Other Ingredients

Acrylate Copolymer, Aloe Barbadensis Leaf (Aloe Vera Gel), Juice, Aqua (Deionized Water), Butyloctyl Salicylate, Camellia Sinensis (Green Tea), Extract, Caprylic/Capric Triglyceride, Cetearyl Oliviate, Cetyl Alcohol, Citric Acid, Ethylhexylglycerin, Glitter: Regenerated Cellulose/Rayon, FD&C Yellow #5, D&C Red 34, Aluminium; Glycerin, Glyceryl Stearate, Helianthus Annuus (Sunflower) Oil, Parfum (Fragrance), Phenoxyethanol, Polysorbate-20, Sorbitan Oliviate, Styrene/Acrylates Copolymer, Tocopherol Acetate (Vitamin E), Zemea (Corn) Propanediol.

Other Information

Protect this product from excessive heat and direct sunlight.

Questions or comments?

Contact us directly at **1-888-601-0441**

Package Labeling:

ONLY THE BEST.

Drug Facts		Purpose
Active Ingredients		Sunscreen
Homosalate	10.00%	Sunscreen
Octocrylene	5.00%	Sunscreen
Octisalate	5.00%	Sunscreen
Zinc Oxide	4.00%	Sunscreen

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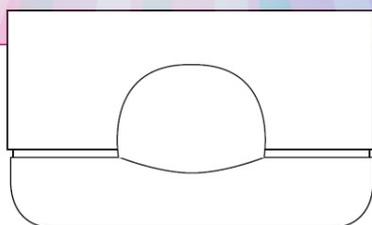
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Questions or Comments?
Contact us directly at 1-888-601-0441
Dist. by: par. Sunshine & Glitter
NJ Products Association LLC
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www.sunshineglitter.com
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 Bleed

 Printing area (text, logos, ...)

SEA STAR SPARKLE SPF 50 GLITTER SUNSCREEN COTTON CANDY

homosalate, octocrylene, octisalate, zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72085-308
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	100 mg in 1 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	50 mg in 1 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 g

ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)

ZINC CATION

40 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
REGENERATED CELLULOSE (UNII: BX81F82EWG)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
D&C RED NO. 34 (UNII: BAN556989E)	
ALUMINUM (UNII: CPD4NFA903)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CORN (UNII: 0N8672707O)	
PROPANEDIOL (UNII: 5965N8W85T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72085-308-00	113 g in 1 PACKET; Type 0: Not a Combination Product	08/10/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	08/10/2020	

Labeler - MJ Products Association, LLC (081022202)