NU SKIN AGELOC ME DAY S1 BROAD SPECTRUM SPF 25- homosalate, octinoxate, avobenzone, and octocrylene lotion NSE Products, Inc.

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.

Nu Skin® ageLOC Me[™] Day S1 Broad Spectrum SPF 25

Drug Facts

Active Ingredient

Homosalate (4.0%), Octinoxate (3.5%), Avobenzone (3.0%), Octocrylene (2.79%)

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
- Stop use and ask a doctor if rash occurs
- **When using this product** keep out of eyes. Rinse with water to remove.
- **Keep out of reach of children.** If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- After applying serum and allowing it to dry, apply moisturizer in an upward, outward motion to the face and neck. If layering additional products, allow each to dry completely before applying the next.
- **Sunprotection 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.—2p.m. Wear long-sleeved shirts, pants, hats, and sunglasses.
- Apply liberally 15 minutes before sun exposure
- Use a water resistant sunscreen if swimming or sweating
- Reapply at least every 2 hours
- Children under 6 months: Ask a doctor

Inactive Ingredients

Water (Aqua), Butylene Glycol, Caprylyl Methicone, Nylon-6/12, Octyldodecyl Neopentanoate, Glycerin, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Oryza Sativa (Rice) Extract, Methoxy PEG/PPG-25/4 Dimethicone, Hydrogenated Phosphatidylcholine, Caprylic/Capric Triglyceride, 4-t-Butylcyclohexanol, Hydroxyacetophenone, Polyisobutene, Schizandra Chinensis Fruit Extract, Camellia Sinensis Leaf Extract, Allantoin, Pentylene Glycol, 7-Dehydrocholesterol, Paeonia Lactiflora Extract, Epilobium Angustifolium Flower/Leaf/Stem Extract, Tocopherol, Narcissus Tazetta Bulb Extract, PEG-7 Trimethylolpropane Coconut Ether, Sorbitan Isostearate, Hydroxyphenyl Propamidobenzoic Acid, Bis-PEG/PPG-20/5 PEG/PPG-20/5 Dimethicone, Zea Mays (Corn) Oil, Xanthan Gum, Aminomethyl Propanol, Disodium EDTA, Chlorphenesin

Other Information

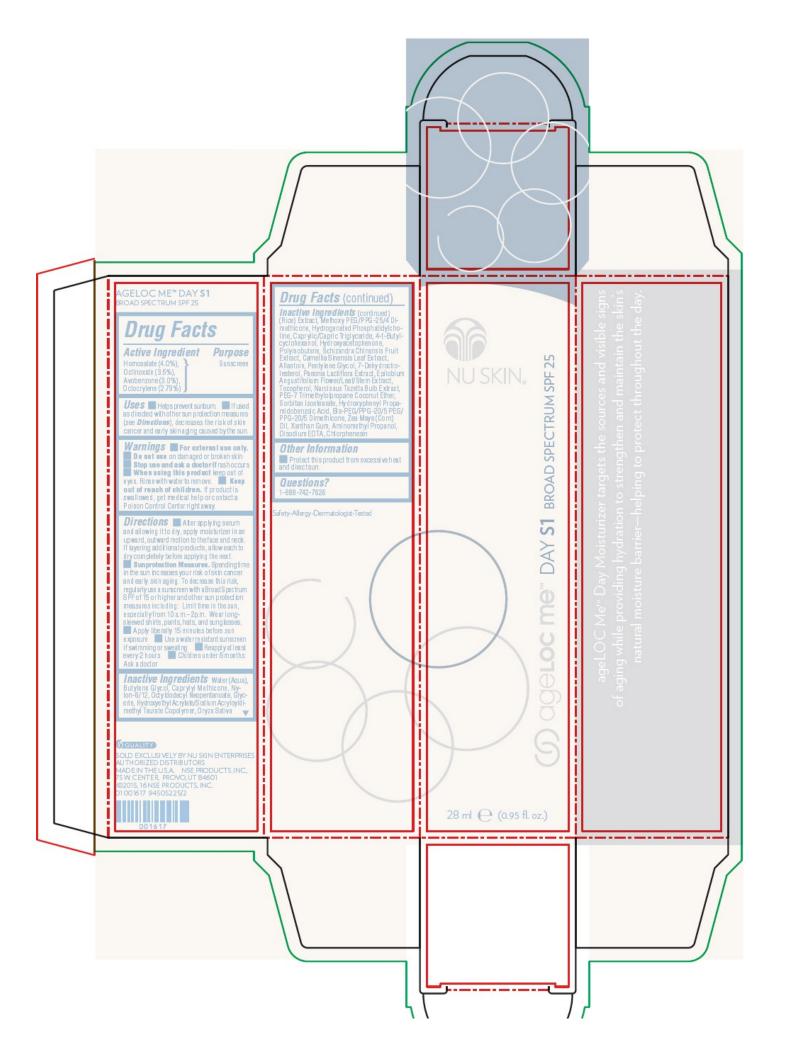
Protect this product from excessive heat and direct sun.

Questions?

1-888-742-7626

PRINCIPAL DISPLAY PANEL - 28 ml Cartridge Carton

NU SKIN $_{\circledR}$ age**LOC me^{\intercal M}** DAY **S1** BROAD SPECTRUM SPF 25 28 ml **e** (0.95 fl . oz.)





NU SKIN AGELOC ME DAY S1 BROAD SPECTRUM SPF 25

homosalate, octinoxate, avobenzone, and octocrylene lotion

Product Information	roduct Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62839-1617			
Route of Administration	TOPICAL					

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
Homosalate (UNII: V06SV4M95S) (Homosalate - UNII:V06SV4M95S)	Ho mo sa late	40 mg in 1 mL		
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	35 mg in 1 mL		
Avobenzone (UNII: G63QQF2NOX) (Avobenzone - UNII:G63QQF2NOX)	Avobenzone	30 mg in 1 mL		
Octocrylene (UNII: 5A68WGF6WM) (Octocrylene - UNII:5A68WGF6WM)	Octocrylene	27.9 mg in 1 mL		

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Butylene Glycol (UNII: 3XUS85K0RA)	
Octyldodecyl Neopentanoate (UNII: X8725R883T)	
Caprylyl Trisiloxane (UNII: Q95M2P1KJL)	
Pentylene Glycol (UNII: 50 C130 7PZG)	
Glycerin (UNII: PDC6A3C0OX)	
Medium-Chain Triglycerides (UNII: C9H2L21V7U)	
Rice Germ (UNII: 7N2B70SFEZ)	
Xanthan Gum (UNII: TTV12P4NEE)	
4-tert-Butylcyclohexanol (UNII: K0H1405S9C)	
Chlorphenesin (UNII: I670 DAL4SZ)	
Allantoin (UNII: 344S277G0Z)	
Edetate Disodium (UNII: 7FLD91C86K)	
Epilobium Angustifolium Flowering Top (UNII: 08H094218D)	
Green Tea Leaf (UNII: W2ZU1RY8B0)	
Sorbitan Isostearate (UNII: 01S2G2C1E4)	
Hydroxyphenyl Propamidobenzoic Acid (UNII: 25KRT26H77)	
Schisandra Chinensis Fruit (UNII: ABS794681C)	
Corn Oil (UNII: 8470G57WFM)	
Narcissus Tazetta Bulb (UNII: K17762966S)	
7-Dehydrocholesterol (UNII: BK1IU07GKF)	
Aminomethylpropanol (UNII: LU49E6626Q)	

	Packaging					
Ш	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
	NDC:62839-1617-	1 in 1 CARTON	05/01/2016			
	1	28 mL in 1 CARTRIDGE; Type 0: Not a Combination Product				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH FINAL	part352	05/01/2016		

Labeler - NSE Products, Inc. (803486393)

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