

ULTIMATE SHEER SUNSCREEN SPF 85 RITE AID- avobenzene 3.00% homosalate 15.00% octisalate 5.00% octocrylene 4.50% oxybenzone 6.00% lotion

Ride Aid

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.

Active ingredients	Purpose
Avobenzene 3.00%.....	Sunscreen
Homosalate 15.00%.....	Sunscreen
Octisalate 5.00%.....	Sunscreen
Octocrylene 4.50%.....	Sunscreen
Oxybenzone 6.00%.....	Sunscreen

Uses • helps prevent sunburn

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 product is 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
- **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
 - children under 6 months: Ask a doctor

Inactive ingredients Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Acrylates/C12-22 Alkyl Methacrylate Copolymer, Acrylates/ Dimethicone Copolymer, Beeswax, BHT, Butyloctyl Salicylate, Cyclopentasiloxane, Dimethyl Capramide, Dipotassium Glycyrrhizate, Disodium EDTA, Ethylhexylglycerin, Fragrance, Glyceryl Stearate, Methylisothiazolinone, PEG-100 Stearate, Polyester-8, Silica, Styrene/ Acrylates Copolymer, Triethanolamine, Water.

May Also Contain Benzisothiazoline, Chlorphenesin

daylogic™

ultimate sheer

SUNSCREEN LOTION

BROAD SPECTRUM SPF 85

UVA/UVB Protection
Dermatologist Tested
Water Resistant (80 Minutes)
Oil-Free
Light, Clean Feel

85

3 FL OZ (89mL)



Drug Facts

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Other information • protect this product from excessive heat and direct sun • may stain fabrics.

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May Also Contain Benzisothiazoline, Chlorphenesin.



DISTRIBUTED BY: RITE AID
30 HUNTER LANE
CAMP HILL, PA 17011

MADE IN THE USA
WITH US AND
IMPORTED MATERIALS

The Skin Cancer Foundation recommends this product as an effective UV sunscreen.

**look great
feel great**
100% GUARANTEE
or your money back

ULTIMATE SHEER SUNSCREEN SPF 85 RITE AID

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-1111
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Avobenzene (UNII: G63QQF2NOX) (AVOBENZENE - UNII:G63QQF2NOX)	Avobenzene	3.00 g in 100 mL
Homosalate (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	Homosalate	15.00 g in 100 mL
Octisalate (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	Octisalate	5.00 g in 100 mL

Octocrylene (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	Octocrylene	4.50 g in 100 mL
Oxybenzone (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	Oxybenzone	6.00 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
Butyloctyl Salicylate (UNII: 2EH13UN8D3)	
CYCLOMETHICONE 5 (UNII: 0TH5PC10R)	
Dimethyl Capramide (UNII: O29Y6X2JEZ)	
GLYCYRRHIZINATE DIPO TASSIUM (UNII: CA2Y0FE3FX)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
Ethylhexylglycerin (UNII: 147D247K3P)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
Methylisothiazolinone (UNII: 229D0E1QFA)	
PEG-100 Stearate (UNII: YD01N1999R)	
POLYESTER-8 (1400 MW, CYANODIPHENYLPROPENYL CAPPED) (UNII: T9296U138P)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STYRENE/ACRYLAMIDE COPOLYMER (MW 500000) (UNII: 5Z4DPO246A)	
TROLAMINE (UNII: 9O3K93S3TK)	
Water (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-1111-3	89 mL in 1 TUBE; Type 0: Not a Combination Product	01/20/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	01/20/2014	

Labeler - Ride Aid (014578892)

Establishment

Name	Address	ID/FEI	Business Operations
Product Quest Mfg.		927768135	manufacture(11822-1111) , label(11822-1111)

Revised: 10/2017

Ride Aid