# AVOBENZONE, HOMOSALATE, OCTISALATE- avobenzone, homosalate, octisalate lotion DOLLAR GENERAL CORPORATION

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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## Sheer Touch Sunscreen SPF 70 D35.000/D35AA

### **Active Ingredients**

Avobenzone 3%

Homosalate 15%

Octisalate 5%

## **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 a doctor if

rash occurs

## Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control right away.

#### **Directions**

- apply liberaly 15 minutes before sun exposure
- apply to all skin exposed to the sun
- 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 Broad Spectrum SPF value of 15 or higher and other sun protection measures including:
- limit time in the sun, especialy from 10 a.m.-2 p.m.
- wear long-sleeved shirts, pants, hats and sunglasses
- children under 6 months of age: Ask a doctor

#### Other information

protect the product in this container from excessive heat and direct sun

## **Inactive ingredients**

water, butyloctyl salicylate, diethylhexyl 2,6-naphthalate, styrene/acrylates copolymer, silica, glyceryl stearate, PEG-100 stearate, dimethicone, potassium cetyl phosphate, benzyl alcohol, beeswax, caprylyl methicone, aluminum starch octenylsuccinate, cetyl dimethicone, behenyl alcohol, acrylates/C12-22 alkyl methacrylate copolymer, xanthan gum, chlorphenesin, dimethicone/PEG-10/15 crosspolymer, sodium polyacrylate, fragrance, glycerin, disodium EDTA, ethylhexyl stearate, tocopheryl acetate, BHT, trideceth-6, pentylene glycol, benzyl benzoate

#### Claims

This product is not manufacturedor distributed by Johnson & Johnson Corporation., distributor of Neutrogena Ultra Sheer Dry-Touch Sunscreen Broad Spectrum SPF 70.

May stain or damage some fabrics or surfaces.

#### **Adverse Reactions**

100% Satisfaction Guaranteed! (888) 309-9030

**DISTRIBUTED BY** 

OLD EAST MIN CO.

100 MISSION RIDGE,

GOODLETTSVILLE, TN 37072

## Principal display panel

STUDIO SELECTION

SUN

**SHEER TOUCH** 

**SUNSCREEN** 

**SUNSCREEN LOTION** 

**BROAD SPECTRUM SPF 70** 

COMPARE TO NEUTROGENA® ULTRA SHEER® SUNSCREEN Broad Spectrum SPF 70\*

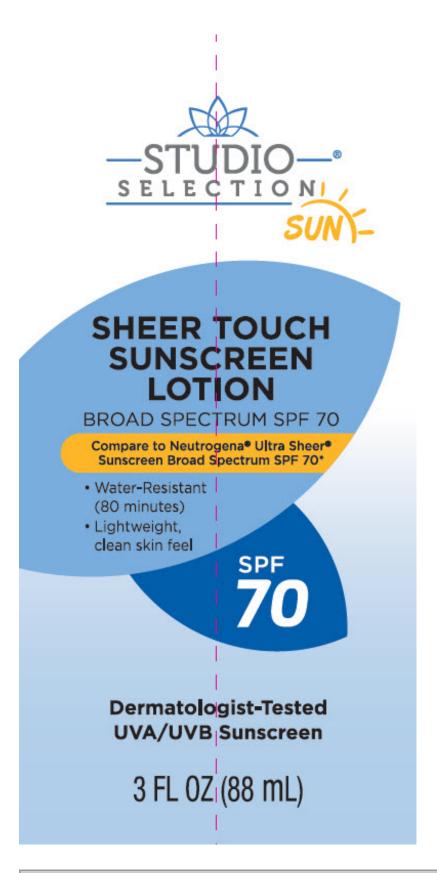
- Water-Resistant (80 Minutes)
- Lightweight, clean skin feel

**SPF 70** 

**Dermatologist-Tested** 

UVA/UVB Sunscreen

3 FL OZ (88 mL)



## **AVOBENZONE, HOMOSALATE, OCTISALATE**

avobenzone, homosalate, octisalate lotion

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75712-935	
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	

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
DIETHYLHEXYL 2,6-NAPHTHALATE (UNII: IODQJ7YGXM)	
STYRENE/ACRYLAMIDE COPOLYMER (500000 MW) (UNII: 5Z4DPO246A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
DIMETHICONE (UNII: 92RU3N3Y10)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: 19PJ0O6294)	
CETYL DIMETHICONE 150 (UNII: 5L694Y0T22)	
DOCOSANOL (UNII: 9G10E216XY)	
BUTYL ACRYLATE/C16-C20 ALKYL METHACRYLATE/METHACRYLIC ACID/METHYL METHACRYLATE COPOLYMER (UNII: 7K68DGG29P)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CHLORPHENESIN (UNII: 1670DAL4SZ)	
DIMETHICONE/PEG-10/15 CROSSPOLYMER (UNII: 21AS8B1BSS)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
BENZYL BENZOATE (UNII: N863NB338G)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75712-935- 21	88 mL in 1 TUBE; Type 0: Not a Combination Product	08/01/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	08/01/2023	

## Labeler - DOLLAR GENERAL CORPORATION (006946172)

## Registrant - Vi-Jon, LLC (088520668)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Vi-Jon, LLC		790752542	manufacture(75712-935)	

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Vi-Jon, LLC		088520668	manufacture(75712-935)	

Revised: 7/2023 DOLLAR GENERAL CORPORATION