DOLLAR GENERAL STUDIO SELECTION SUN SUNSCREEN BROAD SPECTRUM SPF 70- avobenzone, homosalate, octisalate, octocrylene, oxybenzone lotion
Dolgencorp, LLC

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.

Dollar General Studio Selection Sun Sunscreen Lotion Broad Spectrum SPF 70

Active ingredients

Avobenzone 3.0%, Homosalate 15.0%, Octisalate 5.0%, Octocrylene 10.0%, Oxybenzone 6.0%

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 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

• children under 6 months: 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 ingredients

Water, Butylene Glycol, Microcrystalline Cellulose, Glyceryl Stearate, Behenyl Alcohol, Benzyl Alcohol, Diethylhexyl Syringylidenemalonate, Tocopherol (Vitamin E), Retinyl Palmitate (Vitamin A palmitate), Sodium Ascorbyl Phosphate, Stearic Acid, Palmitic Acid, Lauryl Alcohol, Myristyl Alcohol, Cetyl Alcohol, Lecithin, Caprylic/Capric Triglyceride, Chlorphensin, Cellulose Gum, Butylated PVP, Disodium EDTA

Drug Facts

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- may stain or damage some fabrics, materials or surfaces

Inactive ingredients

water, butylene glycol, microcrystalline cellulose, glyceryl stearate, behenyl alcohol, benzyl alcohol, diethylhexyl syringylidenemalonate, tocopherol (vitamin E), retinyl palmitate (vitamin A palmitate), sodium ascorbyl phosphate, stearic acid, palmitic acid, lauryl alcohol, myristyl alcohol, cetyl alcohol, lecithin, caprylic/capric triglyceride, chlorphenesin, cellulose gum, butylated PVP, disodium EDTA

*This product is not manufactured or distributed by Bayer HealthCare LLC, owner of the registered trademarks Coppertone® and Coppertone UltraGuard®.

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STUDIO SELECTION™ SUN

SUNSCREEN LOTION

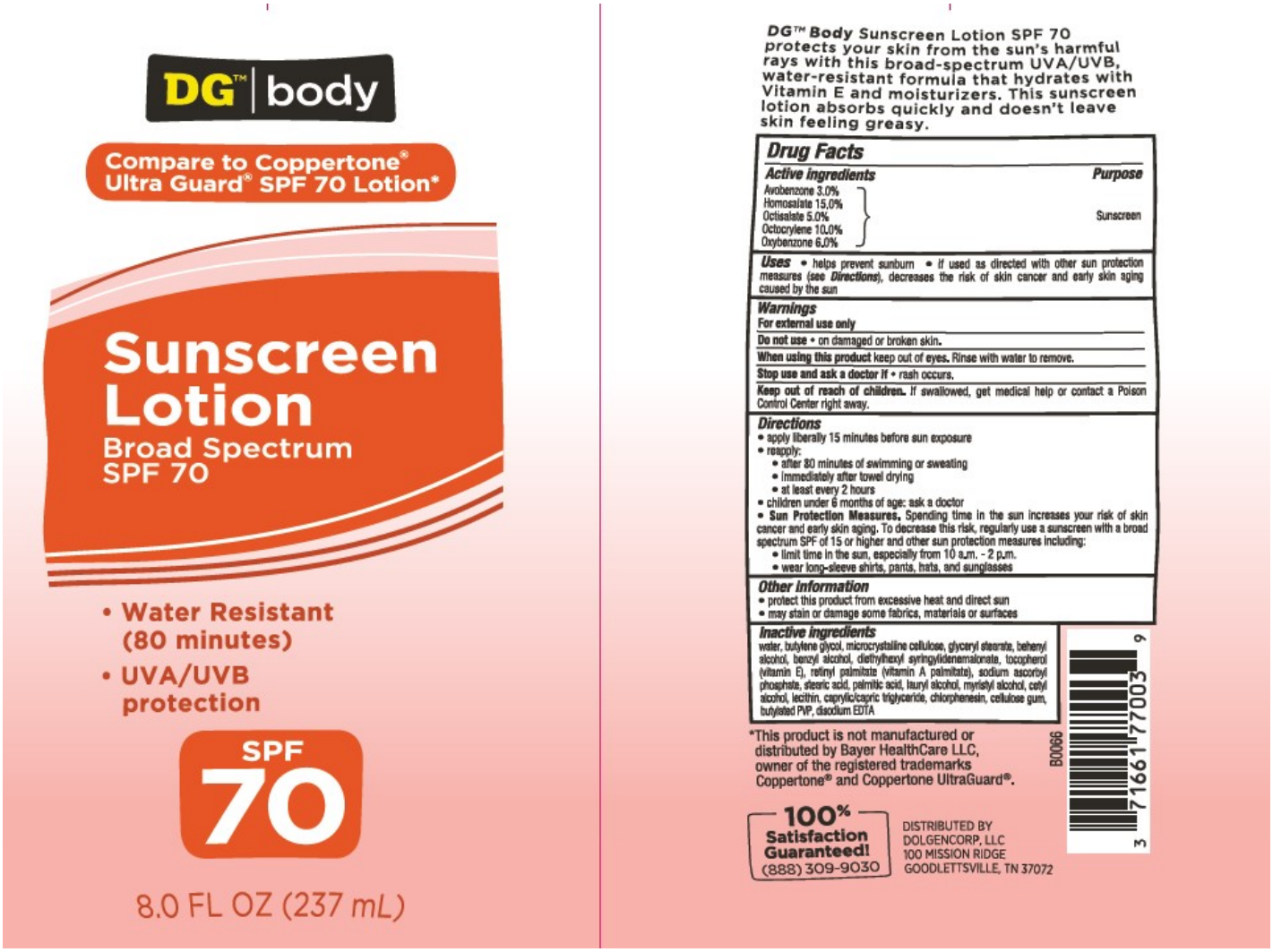
BROAD SPECTRUM SPF 70

Compare to Coppertone® Ultra Guard SPF 70 Lotion*

- Water-Resistant (80 minutes)
- UVA/UVB protection

SPF 70

8 FL OZ (237 mL)



DOLLAR GENERAL STUDIO SELECTION SUN SUNSCREEN BROAD SPECTRUM SPF 70

avobenzone, homosalate, octisalate, octocrylene, oxybenzone lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-036
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	100 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	150 mg in 1 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	60 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
.ALPHA.-TOCOPHEROL, DL- (UNII: 7QWA1RIO01)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
GLYCERYL 1-STEARATE (UNII: 258491E1RZ)	
DOCOSANOL (UNII: 9G1OE216XY)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
DIETHYLHEXYL SYRINGYLIDENEMALONATE (UNII: 3V5U97P248)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
PALMITIC ACID (UNII: 2V16EO95H1)	
LAURYL ALCOHOL (UNII: 178A96NLP2)	
MYRISTYL ALCOHOL (UNII: V42034O9PU)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
EGG PHOSPHOLIPIDS (UNII: 1Z74184RGV)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
SODIUM ASCORBYL PHOSPHATE (UNII: 836SJG51DR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-036-11	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/26/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	10/26/2015	

Labeler - Dolgencorp, LLC (068331990)

Revised: 7/2021 Dolgencorp, LLC