SUNSCREEN- avobenzone, homosalate,otisalate, octocrylene spray
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.

Sunscreen Spray
958

active ingredients

Avobenzone 3%

Homosalate 15%

Octisalate 5%

Octocrylene 10%

Purpose

Sunscreen

uses

- helps prevent sunburn
- if used as directed with other skin protection measures (see Directions), decreases the risk of skin cancer and early skin aging caused by the sun.

Warnings

For external use only

Flammable

Keep away from fire or flame.

- after application, wait until product dries before approaching a souce of heat or flame, or before smoking

Do not use

On damaged or broken skin

When using this product

- Keep out of eyes. Rinse with water to remove.
- contents under pressure. Do not puncture or incinerate. Do not store at temperature above 120

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 Center right away.

directions

- spray liberally and spread evenly by hand 15 minutes before sun exposure.
- apply to all skin exposed to the sun
- do not spray directly into face. Spray on hands then apply to face.
- do not apply in windy conditions
- use in a well-ventilated area and avoid inhalation
- reapply after 80 minutes of swimming or sweating
- immediately after towel drying
- at least every 2 hours
- **Sun protction 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 incluing:
- limit the time in the sun, especially from 10 a.m. - 2 p.m.
- wear long-sleeved shirts, pants, hats and sunglasses
- cildren under 6 month of age: Ask a doctor

Other information

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

inactive ingredients

alcohol denat., acrylates/octylacrylamide copolymer, glycerin, butyloctyl salicylate, triethanolamine, fragrance

claims

May stain or damage some fabrics or surfaces

Oxybenzone & Octinoxate free

disclaimer

This product is not manufactured or distributed by Bayer, distributor of Coppertone Sunscreen Spray Sport Broad Spectrum SPF 70

DSP-TN-15000 DSP-MO-34 SDS-TN-15012

100%

Satisfaction

Guaranteed!

(888)309-9030

DISTRIBUTED BY DOLGENCORP, LLC

100 MISSION RIDGE

GOODLETTSVILLE, TN 37072

958.000/958AA

princpal display panel

STUDIO SELECTION

SUNSCREEN SPRAY

BROAD SPECTRUM SPF 70

Compare to Coppertone Sport Broad Spectrum SPF 70

- Water-resistant (80 minutes
- Continuous spray from any angle

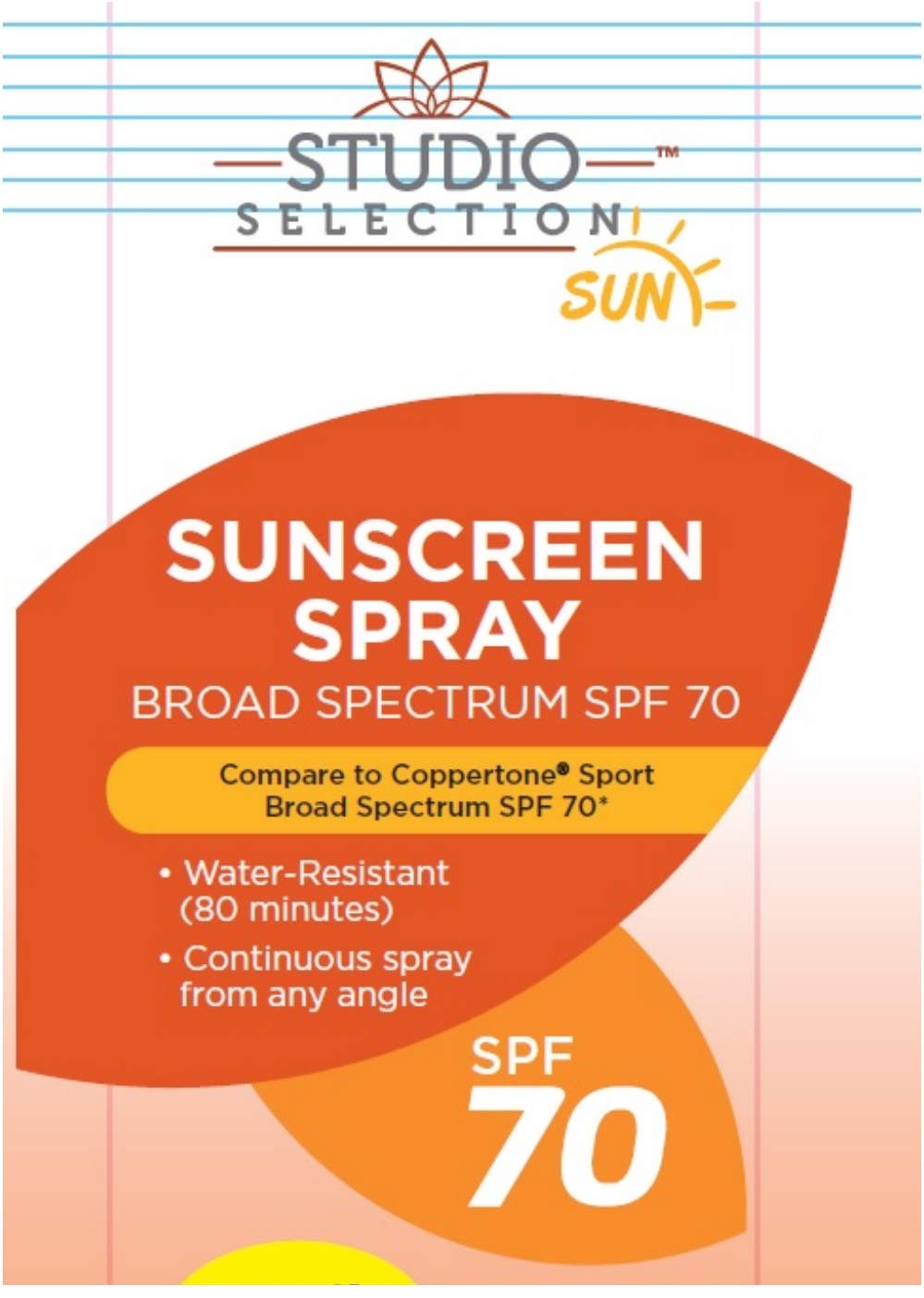
SPF 70

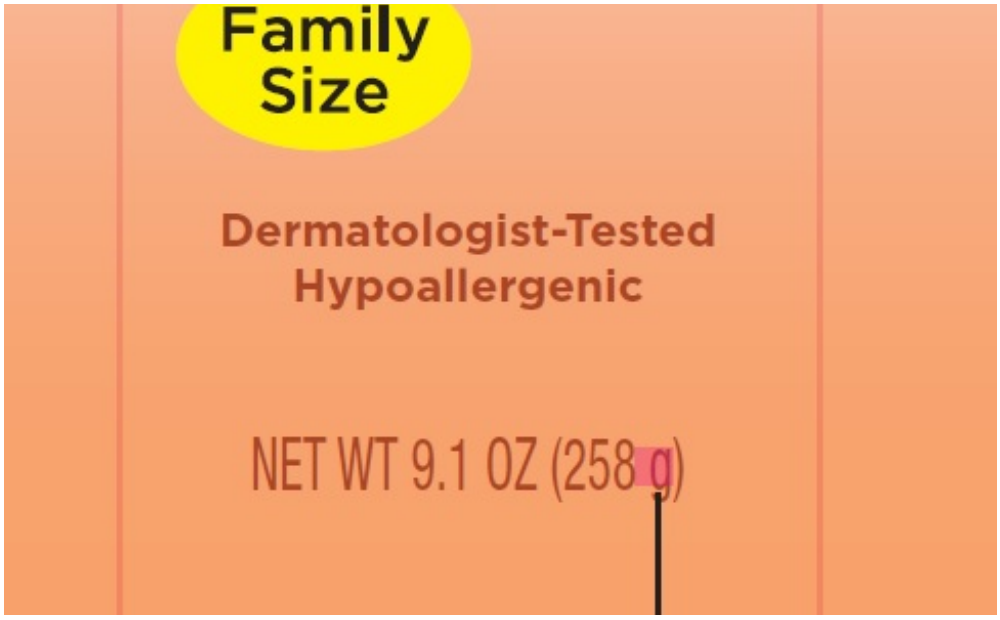
Family Size

Dermatologist-Tested

Hypoallergenic

NET WT 9.1 OZ (258 g)





SUNSCREEN

avobenzone, homosalate,otisalate, octocrylene spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	90 mg in 1 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	225 mg in 1 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	250 mg in 1 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	1000 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
Alcohol (UNII: 3K9958V90M)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:55910-012-15	258 g in 1 CAN; Type 0: Not a Combination Product	10/01/2019	
2	NDC:55910-012-14	156 g in 1 CAN; Type 0: Not a Combination Product	10/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	10/01/2019	

Labeler - DOLGENCORP, LLC (068331990)

Registrant - Vi-Jon, Inc (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, Inc		790752542	manufacture(55910-012)

Revised: 5/2020 DOLGENCORP, LLC