# AVOBENZONE, HOMOSALATE, OCTISALATE- avobenzone, homosalate, octisalate spray Meijer, 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.

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Sunscreen Spray SPF 50 D49.000/D49AA

## **Active Ingredients**

Avobenzone 3%

Homosalate 12%

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

#### **Flammable**

- Flammable: keep away from fire or flame.
- After application, wait until product dries before approaching a source 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°F

## Stop use and ask a doctor if

rash occurs

## Keep out of reach of children.

If swallowed, seek immediate medical attention or call a poison control center.

### **Directions**

- spray liberally and spread evenly by hand 15 minutes before sun exposure
- apply to all skin exposed to the sun
- hold container 4 to 6 inches from the skin to apply
- 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 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
- 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**

alcohol denat., diethylhexyl 2,6-naphthalate, diisopropyl adipate, acrylates/octylacrylamide copolymer, butyloctyl salicylate, neopentyl glycol diheptanoate, fragrance, tocopheryl acetate

### **Claims**

\*This product is not manufactured or distributed by Sun Bum LLC, distributor of Sun Bum® Premium Moisturizing Sunscreen Spray Broad Spectrum SPF 50.

May stain or damage some fabrics or surfaces

### **Adverse Reaction**

Ouestions 1-888-593-0593

**DISTRIBUTED BY:** 

MEIJER DISTRIBUTION, INC.

GRAND RAPIDS, MI 49544

www.meijer.com

**OUR QUALITY GUARANTEE** 

www.meijer.com/satisfaction

# Principal display panel

meijer

sunscreen spray

**BROAD SPECTRUM SPF 50** 

**UVA/UVB SUNSCREEN** 

**SPF 50** 

WATER RESISTANT (80 MINUTES)

FRESH BANANA SCENT

OXYBENZONE, OCTINOXATE AND OCTOCRYLENE FREE

COMPARE TO SUN BUM® PREMIUM MOISTURIZING SUNSCREEN SPRAY\*

NET WT 5.5 OZ (156 g)





# **AVOBENZONE, HOMOSALATE, OCTISALATE**

**TOPICAL** 

avobenzone, homosalate, octisalate spray

**Route of Administration** 

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79481-0949	

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 g	
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	120 mg in 1 g	
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 g	

Inactive Ingredients		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		
DIETHYLHEXYL 2,6-NAPHTHALATE (UNII: I0DQJ7YGXM)		
DIISOPROPYL ADIPATE (UNII: P7E6YFV72X)		
ACRYLATE/ISOBUTYL METHACRYLATE/N-TERT-OCTYLACRYLAMIDE COPOLYMER (40000 MW) (UNII: 7LL6SY9YFV)		
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)		
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79481- 0949-1	156 g in 1 CAN; Type 0: Not a Combination Product	08/08/2023	

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

# **Labeler -** Meijer, Inc. (006959555)

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

Establishment			
Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC.		790752542	manufacture(79481-0949)

Establishment			
Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC.		088520668	manufacture(79481-0949)

Revised: 8/2023 Meijer, Inc.