SUNSCREEN FOR KIDS SPF 50- avebonzone, homosalate, octisalate spray Chain Drug Marketing Association

Quality Choice D42.000/D42AA Sunscreen Spray for Kids SPF 50

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: 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, 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
- 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, acrylates/octylacrylamide copolymer, diisopropyl adipate, neopentyl glycol diheptanoate, butyloctyl salicylate, fragrance, amyl cinnamal, hydroxycitronellal, linalool, tocopheryl acetate

Disclaimer

May stain or damage some fabrics or surfaces

*This product is not manufactured or distributed by Beiersdorf AG, distributor of Coppertone [®]Sunscreen Spray Kids Broad Spectrum SPF 50.

Adverse reaction

100% QC SATISFACTION GUARANTEED

Dstributed by CDMA, Inc.

Novi, MI 48375

www.qualitychoice.com

Questions: 800-935-2362

Principal display panel

NDC 83324-184-05

QC ®

Quality

Choice

*Compare to COPPERTONE ® KIDS

Kids

Sunscreen

Spray

Broad Spectrum SPF 50

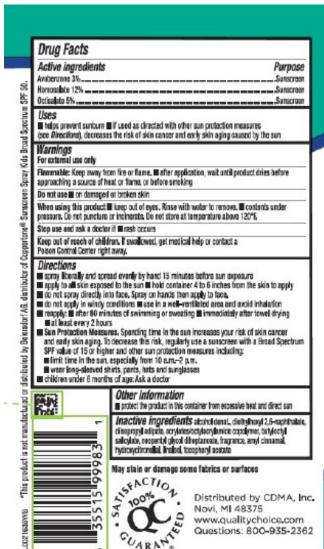
UVA/UVB Sunscreen

Water Resistant (80 Minutes)

Continuous Spray From Any Angle

SPF 50

NET WT 5.5 oz (156 g)







NDC 83324-184-05

Kids Sunscreen **Spray**

Broad Spectrum SPF 50

UVA/UVB Sunscreen

Water Resistant (80 Minutes) Continuous Spray From Any Angle





NET WT **5.5** oz (156 g)

SUNSCREEN FOR KIDS SPF 50

Questions: 800-935-2362

avebonzone, homosalate, octisalate spray

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:83324-184 **Route of Administration TOPICAL**

Active Ingredient/Active Moiety Basis of Strength Ingredient Name 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

II	Inactive Ingredients	
Ш	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)	
.ALPHAAMYLCINNAMALDEHYDE (UNII: WC51CA3418)	
HYDROXYCITRONELLAL (UNII: 8SQ0VA4YUR)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

F	Packaging						
#	# Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:83324-184- 05	156 g in 1 CAN; Type 0: Not a Combination Product	07/08/2024				

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC Monograph Drug	M020	07/08/2024					

Labeler - Chain Drug Marketing Association (011920774)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment					
Name	Address	ID/FEI	Business Operations		
Consumer Product Partners, LLC		119091514	manufacture(83324-184)		

Revised: 3/2025 Chain Drug Marketing Association