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

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

Active ingredient

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

Dstributd by CDMA, Inc.

Novi, MI 48375

www.qualitychoice.com

Questions: 800-935-2362

Principal display panel

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)

SUNSCREEN FOR KIDS SPF 50

avebonzone, homosalate, octisalate spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-991

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	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)	
.ALPHA.-AMYLCINNAMALDEHYDE (UNII: WC51CA3418)	
HYDROXYCITRONELLAL (UNII: 8SQ0VA4YUR)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-991-14	156 g in 1 CAN; Type 0: Not a Combination Product	01/29/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	01/29/2024	

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(63868-991)

Revised: 1/2024 Chain Drug Marketing Association