COPPERTONE GLOW HYDRAGEL SPF 50- avobenzone 3%, homosalate 9%, octisalate 4.5%, octocrylene 9% gel
Beiersdort Inc

Coppertone Glow Hydragel Sunscreen SPF 50

Drug Facts

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Active Ingredients

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Avobenzone 3%, Homosalate 9%, Octisalate 4.5%, Octocrylene 9%

Purpose

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Sunscreen

Uses

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

Warnings

- **For external use only**
- **Flammable.** Keep away from fire or flame.

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 a 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

Directions

■ shake well before each use

■ apply liberally 15 minutes before sun exposure

■ 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-sleeve shirts, pants, hats, and sunglasses

■ children under 6 months: Ask a doctor

Other Information

Other information

■ ***protect this product from excessive heat and direct sun***

■ ***may stain or damage some fabrics or surfaces***

Inactive Ingredients

Inactive ingredients SD alcohol 40-B (57.4% v/v), butyloctyl salicylate, dicaprylyl ether, ethylhexyl isononanoate, PVP, dimethicone/vinyl dimethicone crosspolymer, polyester-27, silica dimethicone silylate, acrylates/C12-22 alkyl methacrylate copolymer, calcium starch octenylsuccinate, fragrance, silica, synthetic fluorphlogopite, iron oxides

Questions

Questions? 1-866-288-3330

Principal Display Panel

Coppertone Sunscreen Gel

Glow Hydragel

SPF 50

Shimmery, Cool & Hydrating

Broad Spectrum SPF 50

Water Resistant (80 Minutes)

Free of oxybenzone, octinoxate, PABA & parabens

4.5 FL OZ (133 mL)

COPPERTONE GLOW HYDRAGEL SPF 50

avobenzone 3%, homosalate 9%, octisalate 4.5%, octocrylene 9% gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66800-4473
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	90 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	90 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	45 mg in 1 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
DICAPRYLYL ETHER (UNII: 77JZM5516Z)	
ETHYLHEXYL ISONONANOATE (UNII: I6KB4GE3K4)	
STARCH, CORN (UNII: O8232NY3SJ)	
ALUMINUM MAGNESIUM HYDROXIDE CARBONATE (UNII: RFJ7QXF345)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (HARD PARTICLE) (UNII: H895X08VNQ)	
POLYESTER-7 (UNII: 0841698D2F)	
BUTYL ACRYLATE/C16-C20 ALKYL METHACRYLATE/METHACRYLIC ACID/METHYL METHACRYLATE COPOLYMER (UNII: 7K68DGG29P)	
FRAGRANCE FRESH CITRUS FLORAL ORC1501495 (UNII: OU4GI2R2WB)	
SILICA DIMETHYL SILYLATE (UNII: EU2PSP0G0W)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
ALCOHOL (UNII: 3K9958V90M)	
OCTENYLSUCCINIC ACID (UNII: 12UZE4X73L)	
MODIFIED CORN STARCH (1-OCTENYL SUCCINIC ANHYDRIDE) (UNII: 461P5CJN6T)	

Product Characteristics

Color	white (white to clear)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66800-4473-4	133 mL in 1 TUBE; Type 0: Not a Combination Product	11/02/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	11/02/2020	

Labeler - Beiersdort Inc (001177906)

Revised: 1/2024 Beiersdort Inc