

SUNSCREEN GLOWING, SPF 30- sunscreen glowing lotion lotion
Cardinal Health, Inc

active ingredient

AVOBENZONE G63QQF2NOX 3.0%
OCTISALATE 4X49Y0596W 5.0%
OCTOCRYLENE 5A68WGF6WM 7.0%
HOMOSALATE V06SV4M95S 9%

purpose

Sunscreen

For external use only Do not use on damaged !i or broken skin.
When using this product keep out of eyes.Rise with water to remove.
Stop use and ask a doctor if rash occurs.
Keep out of reach of children. If swallowed, get medical help or contact a Poison
ControlCenter right away

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Directions

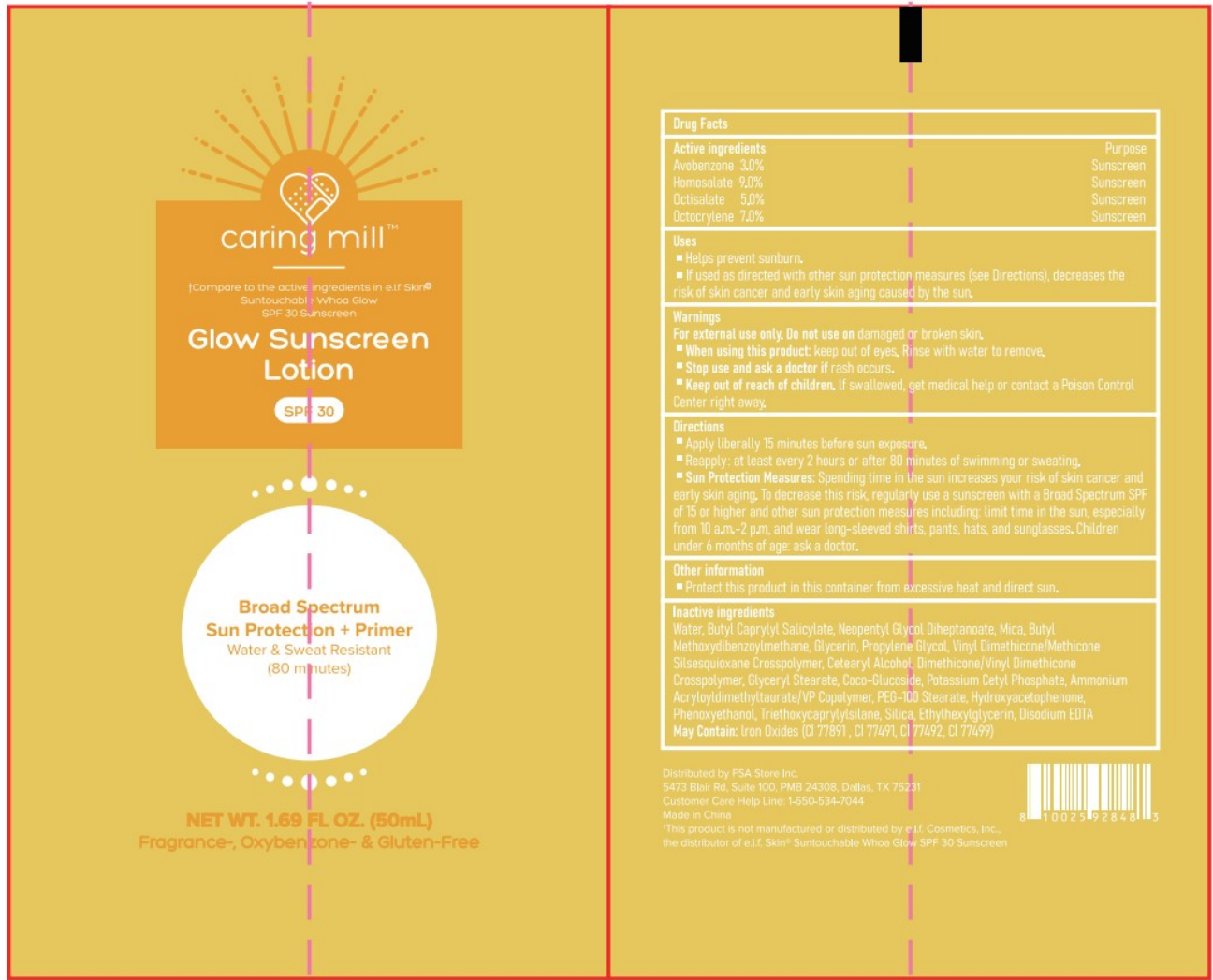
Apply liberally 15 minutes before sun exposure. Reapply: at least every 2 hours or after
80 minutes of swimming or sweating. Sun Protection Measures: \$pending time in the
sun increases your risk of skin cancer andearly skin aging. To decrease this risk,
regularly use a sunscreen with a Broad Spectrum SPfof 15 or higher and other sun
protection measures including: limit time in the sun, especiallyfrom 10 a.m.-2 pm, and
wear long-sleeved shirts, pants, hats, and sunglasses. Childrenunder 6 months of age
ask a doctor.

Apply liberally 15 minutes before sun exposure.
Reapply: at least every 2 hours or after 80 minutes of swimming or sweating.

For external use only. Do not use on damagedor broken skin.

When using this product keep out of eyes. Use with water to remove.
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

WATER
BUTYLOCTYL SALICYLATE
NEOPENTYL GLYCOL DIHEPTANOATE
MICA
CI 77891
BUTYL METHOXYDIBENZOYLMETHANE
GLYCERIN
PROPYLENE GLYCOL
VINYL DIMETHICONE/METHICONE SILSESQUIOXANE CROSSPOLYMER
CETEARYL ALCOHOL
COCO-GLUCOSIDE
GLYCERYL STEARATE
PEG-100 STEARATE
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER
SILICA
POTASSIUM CETYL PHOSPHATE
AMMONIUM ACRYLOYLDIMETHYLTaurate/VP COPOLYMER
CETEARYL ALCOHOL
HYDROXYACETOPHENONE
PHENOXYETHANOL
ETHYLHEXYLGLYCERIN
CI 77492
TRIETHOXYCAPRYLYLSILANE
DISODIUM EDTA
CI 77491
CI 77499



SUNSCREEN GLOWING, SPF 30

sunscreen glowing lotion lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70000-0280
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	4.5 g in 50 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	3.5 g in 50 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	1.5 g in 50 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	2.5 g in 50 mL

Inactive Ingredients

Ingredient Name	Strength
PEG-100 STEARATE (UNII: YD01N1999R)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
CI 77499 (UNII: XM0M87F357)	
COCO GLUCOSIDE (UNII: ICS790225B)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (HARD PARTICLE) (UNII: H895X08VNQ)	
CI 77491 (UNII: 1K09F3G675)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
MICA (UNII: V8A1AW0880)	
CI 77891 (UNII: 15FIX9V2JP)	
CI 77492 (UNII: EX438O2MRT)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
VINYL DIMETHICONE/METHICONE SILSESQUIOXANE CROSSPOLYMER (UNII: 9NH1UDD2RR)	
SILICA (UNII: ETJ7Z6XBU4)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKWBC543X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000-0280-1	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/14/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/14/2024	

Labeler - Cardinal Health, Inc (063997360)