OCEAN POTION FACE WITH CLEAR ZINC OXIDE BROAD SPECTRUM SPF 50 SUNSCREEN- octocrylene, zinc oxide lotion
GNL BRANDS INTERNATIONAL, LLC

Ocean Potion Face with Clear Zinc Oxide Broad Spectrum SPF 50 Sunscreen

Active ingredients

Octocrylene 4.0%, Zinc Oxide 5.0%

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 aging caused by the sun

Warnings

For external use only

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 swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply liberally 15 minutes before sun exposure.
- reapply:
- after 80 minutes of swimming or sweating
- immediately after towel drying

- at least every 2 hours
- children under 6 months of age: Ask a doctor
- **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

Other information

- protect the product in this container from excessive heat and direct sun
- may stain or damage some fabrics, materials or surfaces

Inactive ingredients

water, ethylhexyl stearate, diisopropyl adipate, cetyl dimethicone, polyglyceryl-4 isostearate, cetyl PEG/PPG-10/1 dimethicone, hexyl laurate, phenoxyethanol, beeswax, isohexadecane, hydrogenated castor oil, fragrance, silica, triethoxycaprylylsilane, dimethicone, tocopherol (Vitamin E), tetrasodium EDTA, aloe barbadensis leaf juice, sodium chloride, sodium ascorbyl phosphate.

Label

FRONT

Face • Nose • Lips • Ears

1 FL OZ (30 mL)

Blended and Filled in the U.S.A. with Imported and Domestic Ingredients

8 21039 86077 5

Distributed by
GNL BRANDS INTERNATIONAL LLC
Fort Lauderdale, FL 33306

Drug Facts

Active ingredients **Purpose**
Octocrylene 4.0%, Zinc Oxide 5.0% ..... Sunscreen ▶

web direction
(left side off first)

INSIDE BACK

Drug Facts (continued)

Uses • helps prevent sunburn

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Drug Facts (continued)

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Drug Facts (continued)

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BASE

Drug Facts (continued)

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• limit time in the sun, especially from 10 a.m. – 2 p.m. • wear long-sleeved shirts, pants, hats, and sunglasses ▶

Drug Facts (continued)

Other information • protect the product in this container from excessive heat and direct sun • may stain some fabrics ▶

Drug Facts (continued)

Inactive ingredients aloe barbadensis leaf juice, beeswax, cetyl dimethicone, cetyl PEG/PPG-10/1 dimethicone, diisopropyl adipate, dimethicone, ethylhexyl stearate, fragrance, hexyl laurate, hydrogenated castor oil, isohexadecane, phenoxyethanol, polyglyceryl-4 isostearate, silica, sodium ascorbyl phosphate, sodium chloride, tetrasodium EDTA, tocopherol (vitamin E), triethoxycaprylylsilane, water.

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FACE
with CLEAR ZINC OXIDE
Broad Spectrum SPF 50
SUNSCREEN
1 FL OZ (30mL)

Face · Nose · Lips · Ears

GOES ON CLEAR
MATTE FINISH
WON'T RUN INTO EYES

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Drug Facts (continued)

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OCEAN POTION FACE WITH CLEAR ZINC OXIDE BROAD SPECTRUM SPF 50 SUNSCREEN

octocrylene, zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85638-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	50.2125 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	40 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820DPX33S7)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 5) (UNII: 035JKJ76MT)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
HEXYL LAURATE (UNII: 4CG9F9W01Q)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
CETYL DIMETHICONE 25 (UNII: U4AS1BW4ZB)	
SODIUM ASCORBYL PHOSPHATE (UNII: 836SJG51DR)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
DIISOPROPYL ADIPATE (UNII: P7E6YFV72X)	
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
TOCOPHEROL (UNII: R0ZB2556P8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85638-006-42	30 mL in 1 JAR; Type 0: Not a Combination Product	12/17/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/17/2024	

Labeler - GNL BRANDS INTERNATIONAL, LLC (099638248)

Revised: 8/2025 GNL BRANDS INTERNATIONAL, LLC