

**SUNSCREEN 01- avobenzone,homosalate,octocrylene,octisalate cream
Cosmuses Cosmetics (Ningbo) Co., Ltd.**

SPF30 SUNSCREEN LOTION

Drug Facts

Active ingredients

Avobenzone 3%

Homosalate 10%

Octocrylene 10%

Octisalate 5%

Purpose

Sunscreen

Uses

- Helps prevent sunburn
- If used as directed with other sun protection measures (see Directions), decreased the risk of skin cancer and early skin aging caused by 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 product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply liberally 15minutes before sun exposure
- reapply:
- after 80 minutes of swimming or sweating
- immediate after towel drying
- at least every 2 hours
- Sun Protection Measures

Spending time increases your risk of skin cancer and early skin aging. To decreases 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 10a.m.- 2p.m.
- wear long-sleeve shirts, pants, hats and sunglasses
- children under 6 months

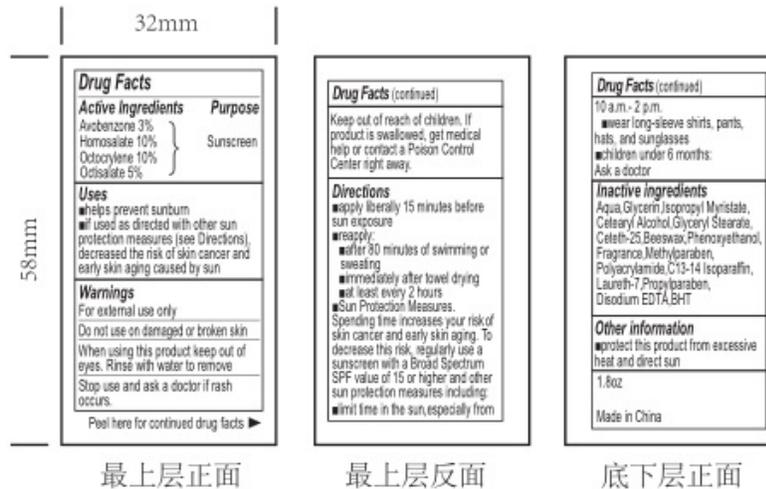
Ask a doctor

Inactive ingredients

Aqua, Glycerin, Isopropyl Myristate, Cetearyl Alcohol, Glyceryl Stearate, Ceteth-25, Beeswax, Phenoxyethanol, Fragrance, Methylparaben, Polyacrylamide, C13-14 Isoparaffin, Laureth-7, Propylparaben, Disodium EDTA, BHT

Other information

Protect this product from excessive heat and direct sun



SUNSCREEN 01

avobenzone,homosalate,octocrylene,octisalate cream

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:82953-017

Route of Administration	TOPICAL
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Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	10 g in 100 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	10 g in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETETH-25 (UNII: 5KLY4IOG20)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
LAURETH-7 (UNII: Z95S6G8201)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
WATER (UNII: 059QF0KO0R)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
POLYACRYLAMIDE (CROSSLINKED; 0.01-0.2 MOLE PERCENT BISACRYLAMIDE) (UNII: RHA9LWJ494)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
FRAGRANCE 13576 (UNII: 5EM498GW35)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82953-017-02	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
2	NDC:82953-017-05	35 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
3	NDC:82953-017-06	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
4	NDC:82953-017-07	120 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
5	NDC:82953-017-03	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
6	NDC:82953-017-04	75 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
7	NDC:82953-017-01	15 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
8	NDC:82953-017-08	240 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
9	NDC:82953-	300 mL in 1 BOTTLE, PLASTIC; Type 0: Not a	05/30/2024	

9	017-09	Combination Product	05/30/2024	
10	NDC:82953-017-10	90 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
11	NDC:82953-017-11	20 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
12	NDC:82953-017-12	45 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
13	NDC:82953-017-13	1.5 mL in 1 POUCH; Type 0: Not a Combination Product	05/30/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	05/30/2024	

Labeler - Cosmuses Cosmetics (Ningbo) Co., Ltd. (725290934)

Registrant - Cosmuses Cosmetics (Ningbo) Co., Ltd. (725290934)

Establishment

Name	Address	ID/FEI	Business Operations
Cosmuses Cosmetics (Ningbo) Co., Ltd.		725290934	manufacture(82953-017)

Revised: 8/2024

Cosmuses Cosmetics (Ningbo) Co., Ltd.