EQUATE SPF 70 ULTRA SUNSCREEN- avobenzone, homosalate, octisalate, octocrylene, oxybenzone lotion Wal-Mart Stores Inc

Walmart Equate SPF 70 Ultra Sunscreen Lotion

Active ingredients

Avobenzone 3.0%, Homosalate 15.0%, Octisalate 5.0%, Octocrylene 10.0%, Oxybenzone 6.0%

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, butylene glycol, microcrystalline cellulose, glyceryl stearate, behenyl alcohol, butylated PVP, benzyl alcohol, palmitic acid, myristyl alcohol, caprylic/capric triglyceride, stearic acid, lauryl alcohol, cetyl alcohol, sodium ascorbyl phosphate, tocopherol (vitamin E), lecithin, cellulose gum, chlorphenesin, disodium EDTA, fragrance.

Label



EQUATE SPF 70 ULTRA SUNSCREEN

avobenzone, homosalate, octisalate, octocrylene, oxybenzone lotion

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-760	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
OXYBENZONE (UNII: 9500S7VE0Y) (OXYBENZONE - UNII:9500S7VE0Y)	OXYBENZ ONE	60 mg in 1 mL	
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	150 mg in 1 mL	
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL	
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	100 mg in 1 mL	
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZ ONE	30 mg in 1 mL	

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
PALMITIC ACID (UNII: 2V16EO95H1)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CHLORPHENESIN (UNII: 1670DAL4SZ)	
N-VINYLPYRROLIDINONE (UNII: 76H9G81541)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
DOCOSANOL (UNII: 9G10E216XY)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
LAURYL ALCOHOL (UNII: 178A96NLP2)	
MYRISTYL ALCOHOL (UNII: V42034O9PU)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K6790BS311)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
TOCOPHEROL (UNII: R0ZB2556P8)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
SODIUM ASCORBYL PHOSPHATE (UNII: 836SJG51DR)	

	Packaging			
# Item Code Package Descr		Package Description	Marketing Start Date	Marketing End Date
	1 NDC:49035-760-	236 mL in 1 TUBE; Type 0: Not a Combination Product	06/04/2019	

Marketing Information			
Marketing Application Number or Monograp Category Citation		Marketing Start Marketing Er Date Date	
OTC Monograph Drug	M020	06/04/2019	

Labeler - Wal-Mart Stores Inc (051957769)

Revised: 10/2024 Wal-Mart Stores Inc