EQUATE BEAUTY ULTRA LIGHT SUNSCREEN SPF70 BROAD SPECTRUMavobenzone, homosalate, octisalate, octocrylene, oxybenzone lotion WALMART STORES INC.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Equate Beauty Ultra Light Sunscreen SPF70 Lotion

Active Ingredient

Avobenzone 3.0%

Homosalate 10.0%

Octisalate 3.0%

Octocrylene 7.0%

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, styrene/acrylates copolymer, dimethicone, acrylates/C12-22 alkyl methacrylate copolymer, propylene glycol, xantham gum, caprylyl methicone, polysilicone-15, cetyl dimethicone, glyceryl stearate, PEG-100 stearate, beeswax, silica, dipotassium glycyrrhizate, trimethylsiloxysilicate, sodium polyacrylate, ethylhexyl stearate, trideceth-6, phenoxyethanol, caprylyl glycol, ethylhexylglycerin, disodium EDTA, fragrance

Label



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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-629
Route of Administration	TOPICAL		

Active ingreal	ent/Active Moiety			
	Ingredient Name	Basis of Streng	gth Strength	
VOBENZONE (UNI	I: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2N	OX) AVOBENZONE	30 mg in 1 mL	
IOMOSALATE (UNI	I: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95	S) HOMOSALATE	100 mg in 1 m	
CTISALATE (UNII:	ISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W) OC		30 mg in 1 mL	
OCTOCRYLENE (UN	ENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM) OCTOCRYLENE		70 mg in 1 mL	
DXYBENZONE (UNI	XYBENZONE (UNII: 9500S7VE0Y) (OXYBENZONE - UNII:9500S7VE0Y) OXYBENZONE		60 mg in 1 mL	
nactive Ingre				
	Ingredient Name		Strengt	
DIMETHICONE (UN	,			
NATER (UNII: 059Q	·			
BUTYL METHACRY UNII: V5RS026Q0H)	LATE/METHYL METHACRYLATE/METHACRYLIC	CACID/STYRENE CROSSPO	DLYMER	
BUTYL ACRYLATE/ COPOLYMER (UNII:	C16-C20 ALKYL METHACRYLATE/METHACRYL 7K68DGG29P)	IC ACID/METHYL METHAC	RYLATE	
PROPYLENE GLYCO	DL (UNII: 6DC9Q167V3)			
XANTHAN GUM (UNII: TTV12P4NEE)				
CAPRYLYL TRISILO	XANE (UNII: Q95M2P1KJL)			
OLYSILICONE-15	(UNII: F8DRP5BB29)			
ETYL DIMETHICO	NE 45 (UNII: IK315POC44)			
GLYCERYL STEARA	TE/PEG-100 STEARATE (UNII: RD25J5V947)			
	UNII: ETJ7Z6XBU4)			
GLYCYRRHIZINATE	DIPOTASSIUM (UNII: CA2Y0FE3FX)			
RIMETHYLSILOXY	SILICATE (M/Q 0.8-1.0) (UNII: 25LXE464L2)			
SODIUM POLYACR	YLATE (8000 MW) (UNII: 285CYO341L)			
	RATE (UNII: EG3PA2K3K5)			
TRIDECETH-6 (UNII	: 3T5PCR2H0C)			
	L (UNII: HIE492ZZ3T)			
	_ (UNII: 00YIU5438U)			
THYLHEXYLGLYC	ERIN (UNII: 147D247K3P)			
EDETATE DISODIU	M (UNII: 7FLD91C86K)			
WHITE WAX (UNII:	7G1J5DA97F)			
Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:49035-629- 09	89 mL in 1 TUBE; Type 0: Not a Combination Product	11/26/2013		
Marketing I	Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	

11/26/2013

OTC monograph not final

part352

Revised: 12/2022

WALMART STORES INC.