EQUATE ZIINC SUNSCREEN ULTRA BROAD SPECTRUM- octinoxate, titanium dioxide, zinc oxide cream
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 Zinc Sunscreen ULTRA Broad Spectrum

Active ingredients

Octinoxate 4.0%

Titanium Dioxide 5.0%

Zinc Oxide 9.1%

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 skin 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 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-sleeve shirts, pants, hats, and sunglasses

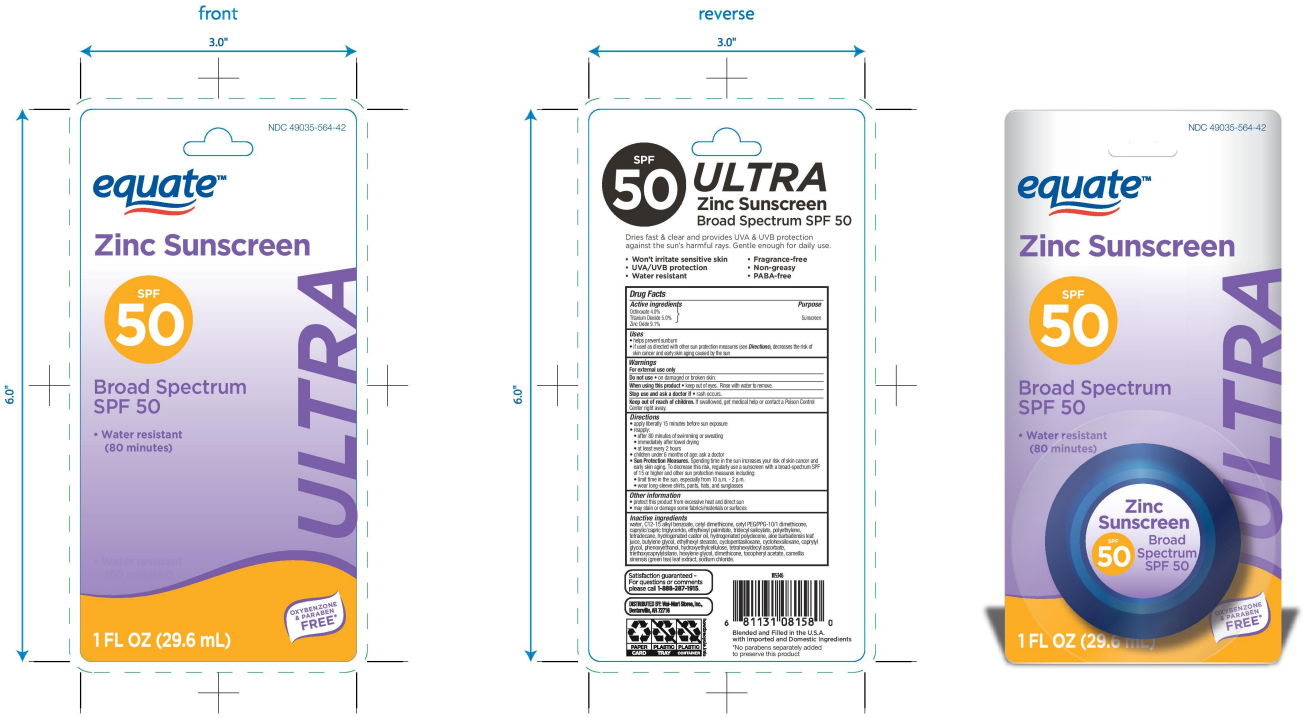
Other information

• protect this product from excessive heat and direct sun

• may stain or damage some fabrics/materials or surfaces

Inactiveingredients

water, C12-15 alkyl benzoate, cetyl dimethicone, cetyl PEG/PPG-10/1 dimethicone, caprylic/capric triglyceride, ethylhexyl palmitate, tridecyl salicylate, polyethylene, tetradecane, hydrogenated castor oil, hydrogenated polydecene, aloe barbadensis leaf juice, butylene glycol, ethylhexyl stearate, cyclopentasiloxane, cyclohexsiloxane, caprylyl glycol, phenoxyethanol, hydroxyethylcellulose, tetrahexyldecyl ascorbate, triethoxycaprylylsilane, hexylene glycol, dimethicone, tocopheryl acetate, camellia sinensis (green tea) leaf extract, sodium chloride

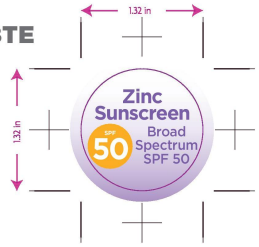


EQ08158E 6.6291″

.7174″

wrap of jar

EQ08158TE



Top of Jar

EQUATE ZIINC SUNSCREEN ULTRA BROAD SPECTRUM

octinoxate, titanium dioxide, zinc oxide cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-564
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	40 mg in 1 mL
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	50 mg in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	91 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
CETYL DIMETHICONE 25 (UNII: U4AS1BW4ZB)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 5) (UNII: 035JKJ76MT)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
TRIDECYL SALICYLATE (UNII: AZQ08K38Z1)	
TETRADECANE (UNII: 03LY784Y58)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
HYDROGENATED POLYDECENE (550 MW) (UNII: U333RI6EB7)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
HYDROXYETHYL CELLULOSE (2000 MPA.S AT 1%) (UNII: S38J6RZN16)	
TETRAHEXYLDECYL ASCORBATE (UNII: 9LBV3F07AZ)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-564-43	1 in 1 PACKAGE	02/20/2019	
1	NDC:49035-564-42	29.6 mL in 1 JAR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	02/20/2019	

Labeler - Walmart Stores Inc. (051957769)

Revised: 10/2021 Walmart Stores Inc.