SPF 30 SHEER SUNSCREEN FACE ULTA- avobenzone 2.00% homosalate 15.00% octinoxate 7.50% octisalate 5.00% stick
Ulta

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.

Drug Facts

Active ingredients	**Purpose**
Avobenzone 2.00%	Sunscreen
Octisalate 3.00%	Sunscreen
Octocrylene 7.00%	Sunscreen
Oxybenzone 3.00%	Sunscreen

Uses • helps prevent sunburn

Warnings
For external use only
Flammable: Do not use near heat, flame, or while smoking.
Do not use on damaged or broken skin
When using this product • Keep out of eyes. Rinse eyes with water to remove.
• Keep away from face to avoid breathing it • Do not puncture or incinerate.
Contents under pressure. Do not store at temperatures above 120°F.
Stop use and ask a doctor if rash occurs

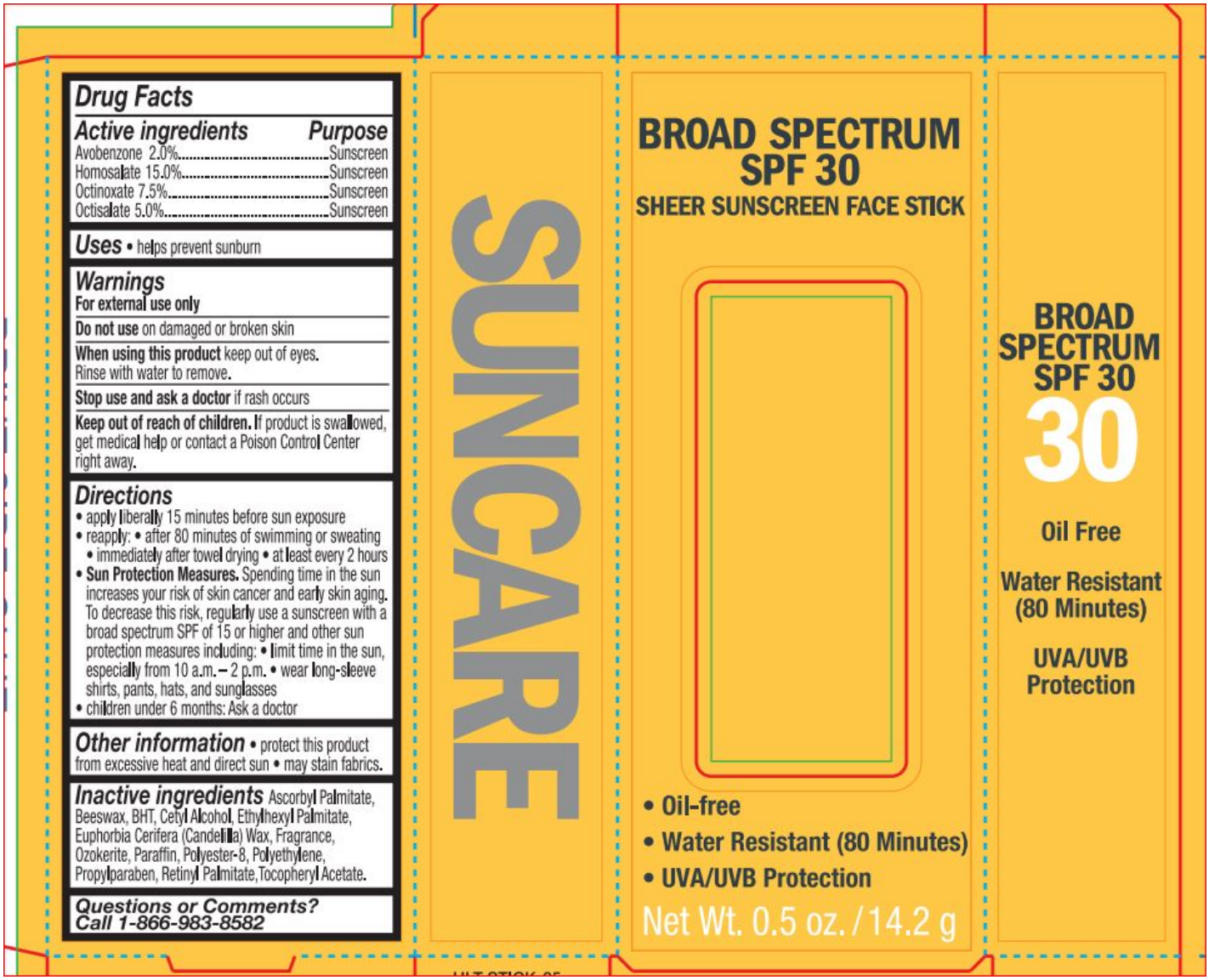
Keep out of reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions
• apply liberally 15 minutes before sun exposure
• reapply:
• after 80 minutes of swimming or sweating
• immediately after towel drying
• at least every 2 hours
• **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
• children under 6 months: Ask a doctor

Inactive ingredients

Ascorbyl Palmitate
Beeswax
BHT
Cetyl Alcohol

Ethylhexyl Palmitate
Euphorbia Cerifera (Candelilla) Wax
Fragrance
Ozokerite
Paraffin
Polyester-8
Polyethylene
Propylparaben
Retinyl Palmitate
Tocopheryl Acetate



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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62296-7778
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Avobenzone (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	Avobenzone	2 g in 100 g
Homosalate (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	Homosalate	15 g in 100 g
Octinoxate (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	Octinoxate	7.5 g in 100 g
Octisalate (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	Octisalate	5 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
Ascorbyl Palmitate (UNII: QN83US2B0N)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
Cetyl Alcohol (UNII: 936JST6JCN)	
Ethylhexyl Palmitate (UNII: 2865993309)	
CANDELILLA WAX (UNII: WL0328HX19)	
CERESIN (UNII: Q1LS2UJO3A)	
PARAFFIN (UNII: I9O0E3H2ZE)	
POLYESTER-8 (1400 MW, CYANODIPHENYLPROPENOYL CAPPED) (UNII: T9296U138P)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	
Propylparaben (UNII: Z8IX2SC1OH)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:62296-7778-1	4.2 g in 1 TUBE; Type 0: Not a Combination Product	01/27/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	01/27/2016	

Labeler - Ulta (608168597)

Registrant - Product Quest Mfg (927768135)

Establishment

Name	Address	ID/FEI	Business Operations
Product Quest Mfg		927768135	manufacture(62296-7778), label(62296-7778)

Revised: 10/2017 Ulta