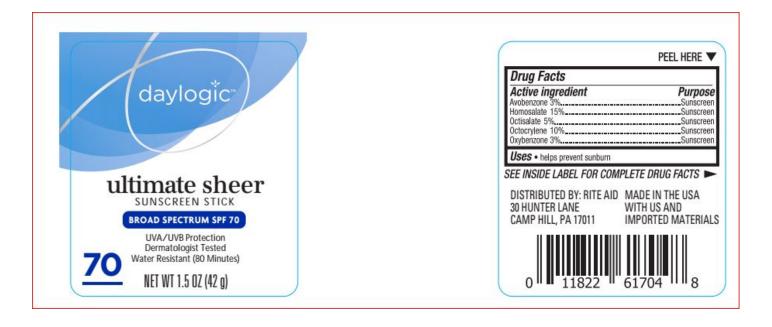
ULTIMATE SHEER SUNSCREEN SPF 70 DAYLOGIC- avobenzone 3% homosalate 15% octisalate 5% octocrylene 10% oxybenzone 3% stick Product Quiest Mfg

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



Drug Facts (continued)

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 Remove cap and twist up tube. See important warnings under "When using this product" • 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,

ADDITIONAL INFORMATION

INSIDE

Drug Facts (continued)

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

Other information

 protect this product from excessive heat and direct sun.
may stain fabrics.

Inactive ingredients

Adipic Acid/Diglycol Crosspolymer, Beeswax, BHT, C12-15 Alkyl Benzoate, Caprylyl Glycol, Fragrance, Neopentyl Glycol Diethylhexanoate, Neopentyl Glycol Diisostearate, Octyldodecyl Neopentanoate, Ozokerite, Paraffin, Petrolatum, Polyester-8, Polyethylene, Silica.

ULTIMATE SHEER SUNSCREEN SPF 70 DAYLOGIC

avobenzone 3% homosalate 15% octisalate 5% octocrylene 10% oxybenzone 3% stick

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64048-1235
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Avobenzone (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	Avobenzone	3 g in 100 g	
Homosalate (UNII: V06SV4M95S) (HOMOSALATE - UNII: V06SV4M95S)	Ho mo salate	15 g in 100 g	
Octisalate (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	Octisalate	5 g in 100 g	
Octocrylene (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	Octocrylene	10 g in 100 g	
Oxybenzone (UNII: 950OS7VE0Y) (OXYBENZONE - UNII:950OS7VE0Y)	Oxybenzone	3 g in 100 g	

Inactive Ingredients	
Ingredient Name	Strength
YELLOW WAX (UNII: 2ZA36H0S2V)	
BUTYLATED HYDRO XYTO LUENE (UNII: 1P9 D0 Z171K)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
Caprylyl Glycol (UNII: 00 YIU5438 U)	
NEOPENTYL GLYCOL DIETHYLHEXANOATE (UNII: U68ZV6W62C)	
Neopentyl Glycol Diisostearate (UNII: 4M6 OQ34JWW)	
Octyldodecyl Neopentanoate (UNII: X8725R883T)	
CERESIN (UNII: Q1LS2UJO3A)	
Paraffin (UNII: 1900E3H2ZE)	
Petrolatum (UNII: 4T6H12BN9U)	
POLYESTER-8 (1400 MW, CYANO DIPHENYLPRO PENO YL CAPPED) (UNII: T9296U138P)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	
SILICON DIO XIDE (UNII: ETJ7Z6XBU4)	

Packaging				
# Item Code Package Description		Marketing Start Date	Marketing End Date	
l	1 NDC:64048-1235-2	43 g in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2016	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part352	12/22/2016		

Labeler - Product Quiest Mfg (927768135)

Registrant - Product Quiest Mfg (927768135)

Establishment

Name	Address	ID/FEI	Business Operations
Product Quiest Mfg		927768135	manufacture(64048-1235)

Revised: 6/2018 Product Quiest Mfg