DR. THROWERS SPF 30 FACE AND BODY SUNSCREEN- avobenzone 3%, homosalate 12%, octisalate 5%, octocrylene 3% lotion
Derma Care Research Labs, LLC

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.

Dr. Thrower's SPF 30 Face and Body Sunscreen Lotion

Active Ingredients

Avobenzone 3%, Homosalate 12%, Octisalate 5%, Octocrylene 3%

Purpose

Sunscreen

Uses

Helps prevent sunburn.

Warnings

For external use only.

Do not use on damaged or broken skin. **When using this product** keep out of eyes. Rinse eyes 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

Sun protection measures: spending time n the sun increases your risk of skin cancer and early skin aging. To decrease risk, regularly use a sunscreen with broad spectrum SPF of 15 or higher and other sun protection measure including:
- limit time in the sun, especially from 10 am to 2 pm
- wear long-sleeved shirts, pants, hats, and sunglasses.

- Children under 6 months: ask a doctor.

Inactive Ingredients

Water, Cetyl Dimethicone, C12-15 Alkyl Benzoate, Hydrated Silica, Styrene/Acrylates Copolymer, Neopentyl Glycol Diheptanoate, Polyester-7, Capryloyl Glycine, Polyester-8, Cinnamomum Zeylanicum Bark Extract, Portulaca Oleracea Extract, Propylene Glycol, Acrylates/Dimethicone Copolymer, BHT, Bisabolol, Caprylyl Glycol, Cedrus Atlantica Bark Extract, Chlorphenesin, Cyclopentasiloxane, Disodium EDTA, Ethylhexyl Stearate, Ethylhexylglycerin, Mannan, Sarcosine, Sodium Polyacrylate, Steareth-2, Steareth-100, Trideceth-6, Xanthan Gum, Phenoxyethanol.

Label

DR. THROWERS SPF 30 FACE AND BODY SUNSCREEN

avobenzone 3%, homosalate 12%, octisalate 5%, octocrylene 3% lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72839-605
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	3 g in 100 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	12 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
STEARETH-100 (UNII: 4OH5W9UM87)	
PURSLANE (UNII: M6S840WXG5)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CINNAMON BARK OIL (UNII: XE54U569EC)	
POLYESTER-8 (1400 MW, CYANODIPHENYLPROPENOYL CAPPED) (UNII: T9296U138P)	
CETYL DIMETHICONE 25 (UNII: U4AS1BW4ZB)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER (UNII: V5RS026Q0H)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	
POLYESTER-7 (UNII: 0841698D2F)	
CAPRYLOYL GLYCINE (UNII: 8TY5YO42NJ)	
WATER (UNII: 059QF0KO0R)	
LEVOMENOL (UNII: 24WE03BX2T)	
PPG-1 TRIDECETH-6 (UNII: 1K7417JX6Q)	
XANTHAN GUM (UNII: TTV12P4NEE)	
2-ETHYLHEXYL ACRYLATE, METHACRYLATE, METHYL METHACRYLATE, OR BUTYL METHACRYLATE/HYDROXYPROPYL DIMETHICONE COPOLYMER (30000-300000 MW) (UNII: S7ZA3CCJ4M)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
SARCOSINE (UNII: Z711V88R5F)	
CEDRUS ATLANTICA BARK (UNII: ITP1Q41UPF)	
STEARETH-2 (UNII: V56DFE46J5)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
YEAST MANNAN (UNII: 91R887N59P)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72839-605-02	57 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/28/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	09/28/2021	

Labeler - Derma Care Research Labs, LLC (116817470)

Revised: 1/2022

Derma Care Research Labs, LLC