DOVE DERMASERIES DRY SKIN RELIEF REPLENISHING FACE CREAM SPF 15-avobenzone, ensulizole, octisalate, octocrylene cream Conopco, Inc. d/b/a Unilever

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.

Dove DermaSeries Dry Skin Relief Replenishing Face Cream SPF 15

Drug Facts

Active ingredients

Avobenzone 2.0%

Ensulizole 1.5%

Octisalate 5.0%

Octocrylene 1.3%

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 product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Massage gently into face and neck for instant relief and lasting hydration
- Apply liberally to face and neck 15 minutes before sun exposure
- 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
- Reapply at least every 2 hours
- Use a water resistant sunscreen if swimming or sweating

Other information

Protect the product in this container from excessive heat and direct sun

Inactive ingredients

Water (Aqua), Glycerin, Stearic Acid, Glycol Stearate, PEG-100 Stearate, Dimethicone, Glyceryl Stearate, Phenoxyethanol, Carbomer, Caprylyl Glycol, Cetyl Alcohol, Xanthan Gum, Disodium EDTA, BHT, Stearamide AMP, Silk Amino Acids, Ascorbic Acid, Panthenol, Tocopheryl Acetate, Biotin, Niacinamide.

Questions or Comments?

1-800-761-3683

For severely dry, itchy skin tested with dermatologists, fragrance free • non-comedogenic broad spectrum SPF 15 sunscreen

Packaging



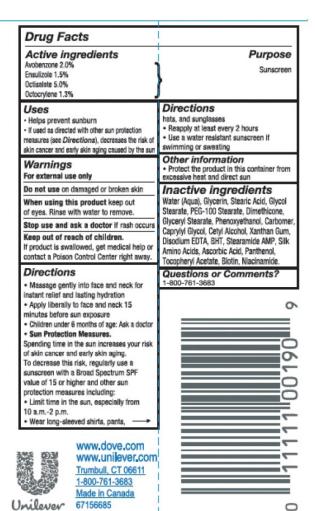
DRY SKIN RELIEF

REPLENISHING

face cream

for severely dry, itchy skin tested with dermatologists fragrance free • non-comedogenic broad spectrum \$PF 15 sunscreen

1.7 FL.0Z. | 50 mL



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avobenzone, ensulizole, octisalate, octocrylene cream

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:64942-1502 Route of Administration TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZ ONE	2 g in 100 mL		
ENSULIZOLE (UNII: 9YQ9DI1W42) (ENSULIZOLE - UNII:9YQ9DI1W42)	ENSULIZOLE	1.5 g in 100 mL		
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL		
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	1.3 g in 100 mL		

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

GLYCERIN (UNII: PDC6A3C0OX)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
GLYCOL STEARATE (UNII: 0324G66D0E)	
PEG-100 STEARATE (UNII: YD01N1999R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
XANTHAN GUM (UNII: TTV12P4NEE)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
STEARAMIDE AMP (UNII: U3K8640346)	
AMINO ACIDS, SILK (UNII: VOLO0EX1IA)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
PANTHENOL (UNII: WV9CM0O67Z)	
.ALPHATOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
BIOTIN (UNII: 6SO6U10H04)	
NIACINAMIDE (UNII: 25X5118RD4)	

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:64942- 1502-1	50 mL in 1 TUBE; Type 0: Not a Combination Product	07/12/2017			
2	NDC:64942- 1502-2	141 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/12/2017			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part352	07/12/2017			

Labeler - Conopco, Inc. d/b/a Unilever (001375088)

Revised: 12/2022 Conopco, Inc. d/b/a Unilever