

**DUKE CANNON STANDARD ISSUE FACE- sunscreen lotion
RNA PHARMA, 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.

EXPIRES 11/22

FACE LOTION

BROAD SPECTRUM
SPF 30

EVERY DAMN DAY

DRUG FACTS

Active Ingredients	Purpose
Avobenzone 3%, Homosalate 6.5%, Octinoxate 7.5%, Octisalate 5%, Oxybenzone 2%	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

• 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

Directions

• Apply generously and evenly 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

Inactive Ingredients

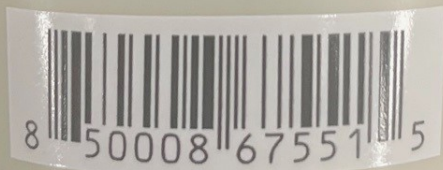
Water, Butyloctyl Salicylate, Glycerin, Silica, Cetearyl Oliviate, Sorbitan Oliviate, Glyceryl Stearate, Ethylhexyl Palmitate, Cetearyl Alcohol, VP/Eicosene Copolymer, Menthyl Lactate, Oryza Sativa (Rice) Bran Extract, Rosmarinus Officinalis (Rosemary) Leaf Extract, Helianthus Annuus (Sunflower) Extract, Tocopherol, Xanthan Gum, Polyacrylate Crosspolymer-6, Caprylyl Glycol, Phenoxyethanol, Ethylhexylglycerin, Disodium EDTA, Fragrance

Other information

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



A PORTION OF PROCEEDS
BENEFITS U.S. VETERANS



Dist. by Duke Cannon Supply Co. | Minneapolis, MN 55401

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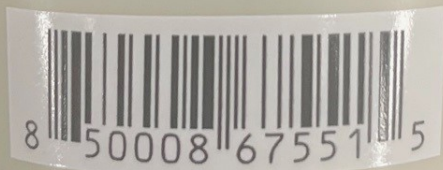
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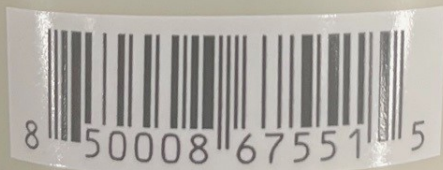
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STANDARD ISSUE

BROAD SPECTRUM
SPF
30

EVERY DAMN DAY
FACE LOTION
2-in-1 Multi-Tasking Moisturizer

UVA + UVB COMPLETE PROTECTION

PROPERTY OF:
DUKE CANNON SUPPLY CO.

MADE IN USA | 1.7 FLOZ (50mL)

DUKE CANNON STANDARD ISSUE FACE

sunscreen lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60717-906
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	0.05 g in 1 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	0.03 g in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	0.065 g in 1 mL
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	0.02 g in 1 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	0.075 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
XANTHAN GUM (UNII: TTV12P4NEE)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
RICE BRAN (UNII: R60QEP13IC)	
ROSEMARY (UNII: IJ67X351P9)	
TOCOPHEROL (UNII: R0ZB2556P8)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
GLYCERYL 1-STEARATE (UNII: 258491E1RZ)	
MENTHYL LACTATE, (-)- (UNII: 2BF9E65L7I)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
EICOSYL POVIDONE (UNII: XQQ9MKE2BJ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60717-906-17	50 mL in 1 TUBE; Type 0: Not a Combination Product	09/21/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	09/21/2021	

Labeler - RNA PHARMA, LLC (079103999)**Establishment**

Name	Address	ID/FEI	Business Operations
RNA PHARMA, LLC		079103999	manufacture(60717-906)

Revised: 9/2021

RNA PHARMA, LLC