EXCHANGE SELECT SUNSCREEN SPF 30- avobenzone homosalate octocrylene oxybenzone lotion
ARMY AND AIR FORCE EXCHANGE SERVICE

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

Avobenzone 2.0 %

Homosalate 10.0%

Octisalate 5.0%

Octocrylene 2.0%

Oxybenzone 2.0%

Purpose

Sunscreen

Uses

- helps prevent sunburn
- higher SPF gives more sunburn protection
- retains SPF after 80 minutes of activity in the water

Warnings

For external use only

When using this product

- keep out of eyes. Rinse with water to remove.

stop use and ask a doctor if

- rash or irritation develops and lasts.

keep out of reach of children

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

Directions

- apply generously before sun exposure and as needed
- children under 6 months of age:ask a doctor
- reapply frequently and after towel drying, swimming or perspiring.

Other Information

- May stain some fabrics
- Sun Alert: Limiting sun exposure, wearing, protective clothing, and using sunscreens may reduce the risks of skin aging, skin cancer, and other harmful effects of the sun.

Inactive Ingredients
Water, Sorbitan Isostearate, Sorbitol, Polyglyceryl-3 Distearate, Triethanolamine, Stearic Acid, VP/Eicosene Copolymer, Benzyl Alcohol, Dimethicone, Barium Sulfate, Carbomer, Simmondsia Chinensis (Jojoba) Seed Oil, Tocopherol, Disodium EDTA, Cocos Nucifera (Coconut) Oil, Mineral Oil, Octadecene/MA Copolymer, Aloe Barbadensis Leaf Extract, Methylparaben, Propylparaben, Fragrance.

Principal Display Panel
exchange
select
X
SUNSCREEN
lotion
SPF
30
Photostable
Broad Spectrum
UVA/UVB Protection
Very Water Resistant
8 FL. OZ. (237mL)


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SUNSCREEN
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ESD4-B
8 FL. OZ. (237 mL)

EXCHANGE SELECT SUNSCREEN SPF 30

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55301-009
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	10 g in 100 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	2 g in 100 g
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	2 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TROLAMINE (UNII: 9O3K93S3TK)	
BARIUM SULFATE (UNII: 25BB7EKE2E)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
TOCOPHEROL (UNII: R0ZB2556P8)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
JOJOBA OIL (UNII: 724GKU717M)	
MINERAL OIL (UNII: T5L8T28FGP)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:55301-009-56	226 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	11/16/2012	

Labeler - ARMY AND AIR FORCE EXCHANGE SERVICE (001695568)

Revised: 11/2012 ARMY AND AIR FORCE EXCHANGE SERVICE