EXCHANGE SELECT SUNSCREEN SPF 45- octinoxate 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 12.0%
Octisalate 5.0%
Octtocrylene 1.6%
Oxybenzone 3.5%

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 the 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 cance, and other harmful effects of the sun.

Inactive ingredients

Water, Sorbitan Isostearate, Sorbitol, Triethanolamine, Octadecene/MA Copolymer, Polyglyceryl-3 Distearate, VP/Eicosene Copolymer, Benzyl Alcohol, Barium Sulfate, Dimethicone, Stearic Acid, Carbomer, Phenethyl Alcohol, Simmondsia Chinensis (Jojoba) Seed Oil, Tocopherol, Disodium EDTA, Aloe Barbadensis Leaf Juice Powder, Methylparaben, Propylparaben.

Principal Display Panel

exchange
select
X
SUNSCREEN
Broad Spectrum Lotion
SPF
45
UVA/UVB Protection
Very Water Resistant


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Broad Spectrum Lotion
SPF
45
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ES07-B
4 FL. OZ. (118 mL)

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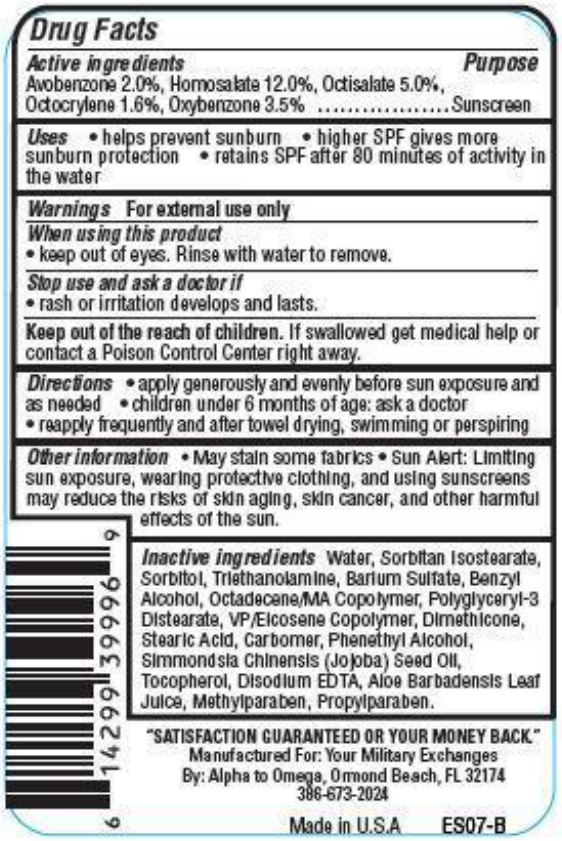

EXCHANGE SELECT SUNSCREEN SPF 45

octinoxate oxybenzone lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55301-001
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	12 g in 100 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	1.6 g in 100 g
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	3.5 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	
SORBITOL (UNII: 506T60A25R)	
BARIUM SULFATE (UNII: 25BB7EKE2E)	
POLYGLYCERYL-3 DIISOSTEARATE (UNII: 46P231IQV8)	

PHENYLETHYL ALCOHOL (UNII: ML9LGA7468)	
TROLAMINE (UNII: 9O3K93S3TK)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
JOJOBA OIL (UNII: 724GKU717M)	
TOCOPHEROL (UNII: R0ZB2556P8)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55301-001-13	113 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	11/15/2012	

Labeler - ARMY AND AIR FORCE EXCHANGE SERVICE (001695568)

Revised: 11/2012 ARMY AND AIR FORCE EXCHANGE SERVICE