# EXCHANGE SELECT SHEER DRY TOUCH SUNSCREEN SPF 70- avobenzone homosalate octisalate 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.

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## **Drug Facts:**

## **Active Ingredients**

Avobenzone 3.0 %

Homosalate 15.0%

Octisalate 5.0%

Octocrylene 2.8%

Oxybenzone 6.0%

## **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

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

• May stain some fabrics

#### **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 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 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-sleeve shirts, pants, hats and sunglasses
- children under 6 months of age: Ask a doctor

#### Other information

• protect this product from excessive heat and direct sun

## **Inactive ingredients**

Water, Styrene/Acrylates Copolymer, Butyloctyl Salicylate, Glyceryl Stearate, Silica, PEG-100 Stearate, Acrylates/C12-22 Alkyl Methacrylate Copolymer, Caprylyl Methicone, Sodium Polyacrylate, Benzyl Alcohol, Beeswax, Ethylhexyl Stearate, Dimethicone, Cetyl Dimethicone, Xanthan Gum, Trimethylsiloxysilicate, Trideceth-6, Disodium EDTA, Dipotassium Glycyrrhizate, BHT, Chlorphenesin, Fragrance.

## **Principal Display Panel**

exchange

select

X

Sheer

**SUNSCREEN** 

70

**BROAD SPECTRUM SPF 70** 

**UVA/UVB PROTECTION** 

Water Resistant

(80 Minutes)

LIGHT, CLEAN FEEL

**DERMATOLOGIST** 

**TESTED** 

oil free

won't clog pores

3 FL.OZ. (88 mL)



### Questions or comments?

Call 1-396-673-2024

## **EXCHANGE SELECT SHEER DRY TOUCH SUNSCREEN SPF 70**

avobenzone homosalate octisalate octocrylene oxybenzone lotion

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55301-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 g		
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	15 g in 100 g		
OCTISALATE (UNII: 4X49 Y0596 W) (OCTISALATE - UNII:4X49 Y0596 W)	OCTISALATE	5 g in 100 g		
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	2.8 g in 100 g		
OXYBENZONE (UNII: 950OS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	6 g in 100 g		

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
BUTYLO CTYL SALICYLATE (UNII: 2EH13UN8D3)	
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)	
GLYCERYL MONOSTEARATE (UNII: 230 OU9 XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
CAPRYLYL TRISILO XANE (UNII: Q95M2P1KJL)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
<b>DIMETHICO NE</b> (UNII: 92RU3N3Y1O)	
XANTHAN GUM (UNII: TTV12P4NEE)	
TETRASILANE (UNII: O19 DXJ0 BL4)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)	
BUTYLATED HYDRO XYTO LUENE (UNII: 1P9 D0 Z171K)	
GLYCYRRHIZINATE DIPO TASSIUM (UNII: CA2Y0 FE3FX)	
CHLORPHENESIN (UNII: 1670 DAL4SZ)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55301-012-22	85 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	10/21/2013		