## SELECT BRAND SUNSCREEN - avobenzone, homosalate, octisalate, octocrylene, oxybenzone lotion Select Brand Distributors

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 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 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**

- shake well
- apply generously and evenly 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 materials

#### **Inactive ingredients**

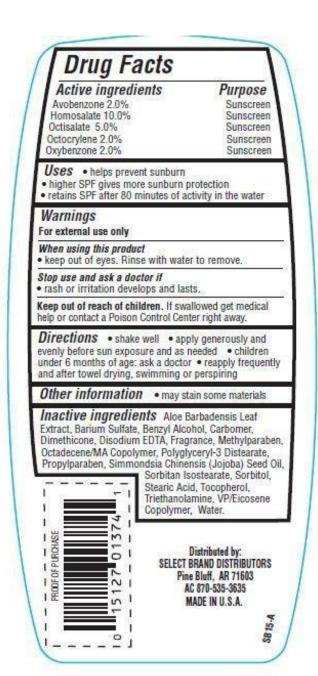
Water, Sorbitan Isostearate, Sorbitol, Polyglyceryl-3 Distearate, Triethanolamine, Stearic Acid, VP/Eicosene Copolymer, Benzyl Alcohol, Dimethicone, Barium Sulfate, Carbomer, Simmondsia Chinensis (Jojoba) Seed Oil, Tocopherol, Disodium EDTA, Octadecene/MA Copolymer, Aloe

Barbadensis Leaf Extract, Methylparaben, Propylparaben, Fragrance.

## **Principal Display Panel**

select brand the lower price name brand SUNSCREEN LOTION 30 SPF Very Water Resistant UVA/UVB Protection with PARSOL 1789 8 FL OZ(237 mL)





## SELECT BRAND SUNSCREEN

avobenzone, homosalate, octisalate, octocrylene, oxybenzone lotion

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:15127-005	
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: 4X49 Y0596W) (OCTISALATE - UNII:4X49 Y0596W)	OCTISALATE	5 g in 100 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	2 g in 100 g
OXYBENZONE (UNII: 9500S7VE0Y) (OXYBENZONE - UNII:9500S7VE0Y)	OXYBENZONE	2 g in 100 g

Inactive Ingredients		
Ingredient Name	Strength	
BENZYL ALCOHOL (UNII: LKG8494WBH)		
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)		
DIMETHICO NE (UNII: 92RU3N3Y1O)		
METHYLPARABEN (UNII: A2I8 C7HI9 T)		
TOCOPHEROL (UNII: R0ZB2556P8)		
PROPYLPARABEN (UNII: Z8IX2SC1OH)		
EDETATE DISO DIUM (UNII: 7FLD91C86K)		
WATER (UNII: 059QF0KO0R)		
SORBITOL (UNII: 506T60A25R)		
TROLAMINE (UNII: 9O3K93S3TK)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
BARIUM SULFATE (UNII: 25BB7EKE2E)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
JOJOBA OIL (UNII: 724GKU717M)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	<b>Marketing End Date</b>
1	NDC:15127-005-16	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	11/21/2012		

# Labeler - Select Brand Distributors (043562370)

Revised: 11/2012 Select Brand Distributors