

**SPF15 SUNSCREEN- avobenzene, homosalate, octocrylene, octisalate stick**  
**Zhejiang Ayan Biotech Co.,Ltd.**

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**SPF15 Sunscreen**

***Drug Facts***

***Active Ingredients***

Homosalate 8% Octocrylene 5.5% Avobenzene 2% Octisalate 2%

***Purpose***

Sunscreen

***Uses***

- Helps prevent sunburn

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

***Directions***

- Apply liberally 15 minutes before sun exposure.
- Use a water resistant sunscreen if swimming or sweating.
- Reapply at least every 2 hours.
- Children under 6 months: Ask a doctor.

### **(Sunscreen) Inactive Ingredients**

Mineral Oil, Beeswax, Ozokerite, Cera Microcristallina, Cocos Nucifera (Coconut) Oil, Ethylhexyl Palmitate, Ethylhexyl Palmitate, Polyisobutene, Polyethylene, Tridecyl Trimellitate, Butyrospermum Parkii (Shea Butter), BHT, Tocopheryl Acetate, Flavor. May Contain: Blue 1 Lake (CI 42090), Yellow 5 Lake (CI 19140), Yellow 6 Lake (CI 15985), Titanium Dioxide (CI 77891), Red 6 Lake (CI 15850), Red 7 Lake (CI 15850), Red 27 Lake (CI 45410), Red 28 Lake (CI 45410), Red 30 Lake (CI 73360), Iron Oxides (CI 77492, CI 77491, CI 77499).

### **Other Information**

- Protect from excessive heat and direct sun.

### **Package Labeling:**

**Broad Spectrum SPF 15**  
Sunscreen - Net Wt. 2g

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### ***Inactive Ingredients*** (Sunscreen)

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### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:70412-212
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>AVOBENZONE</b> (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	20 mg in 1 g
<b>HOMOSALATE</b> (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	80 mg in 1 g
<b>OCTOCRYLENE</b> (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	55 mg in 1 g
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	20 mg in 1 g

### Inactive Ingredients

Ingredient Name	Strength
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>YELLOW WAX</b> (UNII: 2ZA36H0S2V)	
<b>COCONUT OIL</b> (UNII: Q9L0O73W7L)	
<b>ETHYLHEXYL PALMITATE</b> (UNII: 2865993309)	
<b>HIGH DENSITY POLYETHYLENE</b> (UNII: UG00KM4WR7)	
<b>TRIDECYL TRIMELLITATE</b> (UNII: FY36J270ES)	
<b>SHEA BUTTER</b> (UNII: K49155WL9Y)	
<b>BUTYLATED HYDROXYTOLUENE</b> (UNII: 1P9D0Z171K)	
<b>METHYL SALICYLATE</b> (UNII: LAV5U5022Y)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70412-212-02	2 g in 1 TUBE; Type 0: Not a Combination Product	10/01/2018	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	10/01/2018	

