

**BULL FROG WATER ARMOR SPORT QUIK 50- avobenzene, homosalate, octisalate, oxybenzone, octocrylene gel**  
**Sun & Skin Care Research, LLC**

*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**

Homosalate: 15%

Octisalate: 5%

Oxybenzone: 6%

Octocrylene: 10%

Avobenzene: 3%

### **Purpose**

Sunscreen

### **Uses**

- helps prevent sunburn
- If used as directed with other skin protection measures (see Directions), decreases the risk of skin cancer and early skin aging caused by the sun.

### **Warnings**

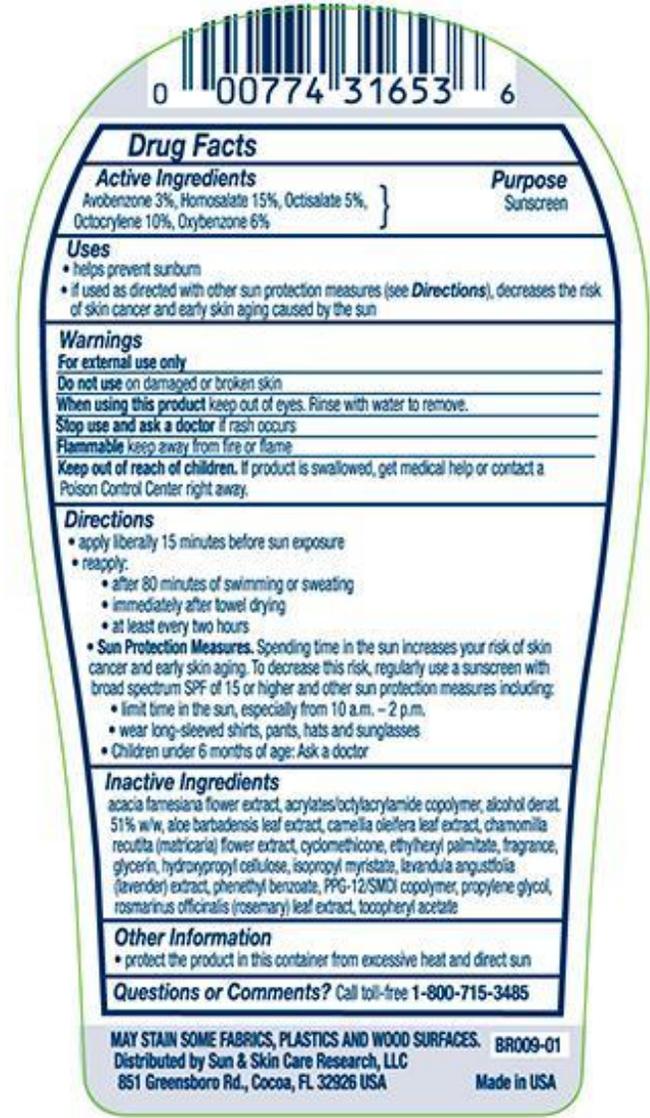
**For external use only. Do not use** on damaged or broken skin. **Stop use and ask a doctor if** rash occurs. **When using this product** keep out of eyes. Rinse with water to remove. **Keep out of the reach of children.** If swallowed, get medical help or contact a Poison Control Center right away. **Flammable** Keep away from fire or flame.

### **Directions**

- Apply liberally 15 minutes before sun exposure
- reapply after 80 minutes of swimming or sweating and 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 value of 15 or higher and other sun protection measures including limiting time in the sun, especially from 10a.m.-2p.m., wear long-sleeved shirts, pants hats and sunglasses.
- children under 6 months: Ask a doctor.

### **Other Information**

- For use on skin only
- Avoid contact with fabric
- Protect this product from excessive heat and direct sun



## BULL FROG WATER ARMOR SPORT QUIK 50

avobenzone, homosalate, octisalate, oxybenzone, octocrylene gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62802-653
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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
OXYBENZONE (UNII: 95OOS7VE0 Y) (OXYBENZONE - UNII:95OOS7VE0 Y)	OXYBENZONE	6 g in 100 g
OCTOCRYLENE (UNII: 5A68 WGF6 WM) (OCTOCRYLENE - UNII:5A68 WGF6 WM)	OCTOCRYLENE	10 g in 100 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
CYCLOMETHICONE (UNII: NMQ347994Z)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
CAMELLIA OLEIFERA LEAF (UNII: 5077EL0C60)	
CHAMOMILE FLOWER OIL (UNII: 60F80Z61A9)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
LAVANDULA ANGUSTIFOLIA FLOWER (UNII: 19AH1RAF4M)	
ROSMARINUS OFFICINALIS FLOWER (UNII: NR1A27F29O)	
HYDROXYPROPYL CELLULOSE (UNII: RFW2ET671P)	
PHENETHYL BENZOATE (UNII: 0C143929GK)	
PPG-12/SMDI COPOLYMER (UNII: 1BK9DDD24E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62802-653-05	147 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	01/01/2014	

**Labeler** - Sun & Skin Care Research, LLC (849772207)

## Establishment

Name	Address	ID/FEI	Business Operations
Sun & Skin Care Research, LLC		849772207	manufacture(62802-653)

Revised: 3/2014

Sun & Skin Care Research, LLC