

**ORLASKIN HYDRATING SUNSCREEN- orlaskin hydrating sunscreen spray  
Guangzhou Fantasy Biotechnology Co.,Ltd**

*Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.*

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**83447-003**

Homosalate 10%

Octocrylene 8%

Octisalate 5%

Avobenzone 3%

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 ageing caused by the sun.

**Warnings**

For external use only.

Avoid contact with eyes , wear protective clothing , hats and eyewear when exposed to the sun . Avoid prolonged exposure in the sun.

Keep out of reach of children.

Do not use on damaged or brokenskin.

Store in a cool , dry place and avoiddirect sunlight.

If using with other sun protection products , consult with a doctor before use.

Do not use on damaged or broken skin

Keep out of reach of children

**Directions**

Shake the spray bottle well before use.

Hold the spray bottle at an appropriate distance from the face and neck . Spray evenly onto the entire face and neck area.

Avoid direct contact with eves . In case of eye contact , rinse immediately with water.

For continued sun protection , reapply every 2 hours

## Other information

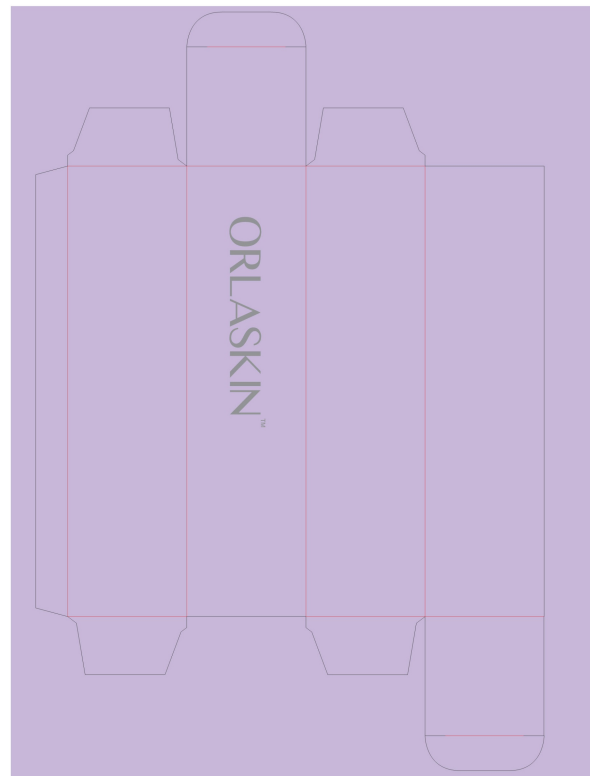
Protect the product in this container from excessive heat and direct sun

## Inactive Ingredients

Auqa , Methacrylate Copolymer , Alcohol Denat , Bisabolol , Hvaluronic Acid, D-alpha-tocopheryl acetate , Terminalia Ferdinandiana , Butyloctyl Salicylate, Sebacic Acid Copolymer , Dicaprylyl Ether , Diheptyl Succinate , Diisooctyl Succinate , Ethyl Ferulat , Isononyl sononanoate , Pelargonium Graveolens Flower Oil

尺寸: 41\*41\*155mm+内托

灰色部分做镭射



## ORLASKIN HYDRATING SUNSCREEN

orlaskin hydrating sunscreen spray

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:83447-003
<b>Route of Administration</b>	CUTANEOUS		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>OCTOCRYLENE</b> (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	8 g in 100 mL
<b>HOMOSALATE</b> (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	10 g in 100 mL
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL

**AVOBENZONE** (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX) | AVOBENZONE | 3 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>HYALURONIC ACID</b> (UNII: S270N0TRQY)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>KAKADU PLUM</b> (UNII: 0ZQ1D2FDLI)	
<b>BUTYLOCTYL SALICYLATE</b> (UNII: 2EH13UN8D3)	
<b>CAPRYLOYL GLYCERIN/SEBACIC ACID COPOLYMER (2000 MPA.S)</b> (UNII: N7YC58165T)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ETHYL ACRYLATE AND METHYL METHACRYLATE COPOLYMER (2:1; 750000 MW)</b> (UNII: P2OM2Q86BI)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>LEVOMENOL</b> (UNII: 24WE03BX2T)	
<b>DICAPRYLYL ETHER</b> (UNII: 77JZM5516Z)	
<b>DIHEPTYL SUCCINATE</b> (UNII: 057N7SS26Y)	
<b>ETHYL FERULATE</b> (UNII: 5B8915UELW)	
<b>ISONONYL ISONONANOATE</b> (UNII: S4V5BS6GCX)	
<b>PELARGONIUM GRAVEOLENS FLOWER OIL</b> (UNII: 3K0J1S7QGC)	
<b>DOCUSATE SODIUM</b> (UNII: F05Q2T2JA0)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83447-003-01	100 mL in 1 BOX; Type 0: Not a Combination Product	10/19/2023	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/19/2023	

**Labeler** - Guangzhou Fantasy Biotechnology Co.,Ltd (619047084)

### Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Fantasy Biotechnology Co.,Ltd		619047084	manufacture(83447-003)

Revised: 10/2023

Guangzhou Fantasy Biotechnology Co.,Ltd