

**FAMILY WELLNESS- diphenhydramine hydrochloride, zinc acetate cream**  
**Zhejiang Jingwei Pharmaceutical Co., Ltd.**

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**Drug Facts**

Inactive Ingredients - cetyl alcohol, glycerol stearate,  
laureth-23, methylparaben, mineral oil, petrolatum, propylene  
glycol, propylparaben, stearic acid, stearyl alcohol, water

Active Ingredients

Diphenhydramine hydrochloride 2 %

Zinc acetate 0.1 %

Purpose

---Topical analgesic

---Skin protectant

Uses for temporary relief from pain and itching associated  
with insect bites, sunburn and minor skin irritations. Dries the  
oozing and weeping of poison: -ivy -oak - sumac

Warnings

For external use only

Do not use

-over large areas of the body

-with any other product containing diphenhydramine,  
including those taken orally

When using this product - avoid contact with eyes

Ask a doctor before use

-on chicken pox

-on measles

Stop use and ask a doctor if:

-symptoms last for more than 7 days

-the condition gets worse

-symptoms clear up and then occur again within a few days

Keep out of reach of children. If swallowed, get medical help  
or contact a Poison Control Center right away.

Directions

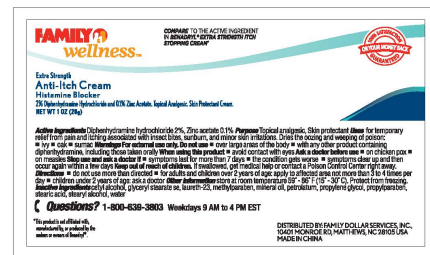
-for adults and children over 2 years of age: apply to affected area

not more than 3 or 4 times per day

-children under 2 years of age: ask a doctor before use

Other information - store at room temperature 59 - 86 F (15 - 30 C)

Questions? 1-800-639-3803 Weekdays 9 AM to 4 PM EST



## FAMILY WELLNESS

diphenhydramine hydrochloride, zinc acetate cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:55621-007
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	2 g in 100 g
<b>ZINC ACETATE</b> (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	0.1 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>LAURETH-23</b> (UNII: N72LMW566G)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>STEARYL ALCOHOL</b> (UNII: 2KR89I4H1Y)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55621-007-01	1 in 1 BOX	11/01/2013	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/01/2013	

**Labeler** - Zhejiang Jingwei Pharmaceutical Co., Ltd. (530876549)

Revised: 10/2023

Zhejiang Jingwei Pharmaceutical Co., Ltd.