

WORMWOOD BALM- chlorhexidine diacetate cream
Shenzhen Furuizhilian keji Co., Ltd.

WORMWOOD BALM

MENTHOL 0.5%

Insect Repellents

Make sure the application area is clean and dry.

Apply to the mosquito bites, or to the limbs or other parts to prevent bites.

For external use only

Don't eat, avoid contact with your eyes.

irritation and redness develop. If condition persists more than 48 hours.

Do not get into eyes.

If contact occurs, rinse eyes thoroughly with water.

If swallowed, get medical help or contact a Poison Control Center right away.

Artemisia Argyi Leaf Extract, Peppermint Extract, Menthol, Beeswax



Wormwood Balm

Drug Facts	
Active ingredient CHLORHEXIDINE DIACETATE 0.2%	Purpose Insect Repellents
Uses ■ For repelling ticks, gnats, biting flies, chiggers and fleas	
Warnings ■ For external use only ■ Don't eat, avoid contact with your eyes ■ Do not use it on damaged skin, otherwise your damaged skin maybe painful	
When using this product ■ Do not get into eyes ■ If contact occurs, rinse eyes thoroughly with water	
Stop use and ask a doctor if ■ Irritation and redness develop. If condition persists more than 48 hours	
Drug Facts(continued)	
Keep out of reach of children	
Directions ■ Make sure the application area is clean and dry ■ Apply to the mosquito bites, or to the limbs or other parts to prevent bites	
Other information Store at 59-86 °F (15-30°C)	
Inactive Ingredients Artemisia Argyi Leaf Extract, Peppermint Extract, Menthol, Beeswax	

WORMWOOD BALM

chlorhexidine diacetate cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84445-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.15 g in 30 g

Inactive Ingredients

Ingredient Name	Strength
CHLORHEXIDINE ACETATE (UNII: 5908ZUF22Y)	
ARTEMISIA ARGYI LEAF (UNII: 2JYC99Q0WZ)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
PEPPERMINT (UNII: V95R5KMY2B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84445-004-01	30 g in 1 BOTTLE; Type 0: Not a Combination Product	07/15/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/15/2024	

Labeler - Shenzhen Furuizhilian keji Co., Ltd. (418598613)**Establishment**

Name	Address	ID/FEI	Business Operations
Shenzhen Furuizhilian keji Co., Ltd.		418598613	manufacture(84445-004)

Revised: 7/2024

Shenzhen Furuizhilian keji Co., Ltd.