ICHTHAMMOL 20%- ichthammol ointment
Chain Drug Marketing Association, Inc.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Quality Choice Ichthammol Ointment - Advanced Drawing Salve
20% Ichthammol - Homeopathic

DRUG FACTS

Active ingredient

Ichthammol 20%

Purpose

Drawing Salve

Keep out of reach of children.

If swallowed, get medical help or contact a poison control center immediately.

Suggested Use

• Treating insect bites and stings from mosquitoes, spider, and bees • Removing splinters and silvers • Treating plant irritations, such as nettles or poison ivy • Treating minor skin infections such as an ingrown toenail • Soothing skin relief • Apply to minor skin irritations

Warnings

• For external use only • Do not use in eyes • Ask a doctor before use if you have deep puncture wounds or serious burns •

Directions

Cleanse affected area. Apply on gauze bandage once or twice a day. Repeat as necessary. Do not rub or massage affected area.

Stop Use and ask a doctor if

Stop use and ask a doctor if redness , irritation, swelling or pain persists.

Inactive Ingredients

Lanolin, Petrolatum

Distributed by CDMA., Inc.©

Novi, MI 48375

www.qualitychoice.com

Questions: 800-935-2362

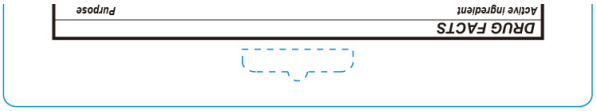
Other Information

Before using any medication, read all label directions

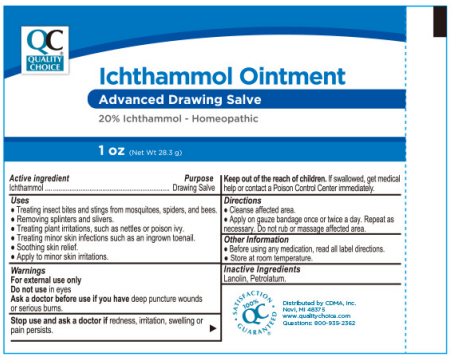
Store at room temperature

Packaging

OUTSIDE BOX



INNER TUBE



ICHTHAMMOL 20%

ichthammol ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-048
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ICHTHAMMOL (UNII: NQ14646378) (ICHTHAMMOL - UNII:NQ14646378)	ICHTHAMMOL	20 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
PETROLATUM (UNII: 4T6H12BN9U)	
LANOLIN (UNII: 7EV65EAW6H)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:83324-048-01	1 in 1 BOX	03/25/2024	
1		28.3 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
unapproved homeopathic		03/19/2024	

Labeler - Chain Drug Marketing Association, Inc. (011920774)

Registrant - Trifecta Pharmaceuticals USA (079424163)

Revised: 7/2024 Chain Drug Marketing Association, Inc.