HYDROPHOR- hydrophor ointment
Akron Pharma Inc.

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.

Hydrophor Ointment

Drug Facts

Active Ingredient:

Petrolatum 42%

Purpose:

Skin protectant

Uses

temporarily protects minor:
- cuts
- scrapes
- burns

temporarily protects and helps relieve
- chapped or cracked skin and lips
- helps protects from the drying effects of wind and cold weather

Warnings

For External Use Only.

When using this product:
- do not get into eyes

Stop use and ask a doctor if
- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few days

Do not use on
- deep or puncture wounds
- animals bites
- serious burns

- **KEEP OUT OF REACH OF CHILDREN.**

Directions

- Apply as needed.

Inactive Ingredient

Ceresin Wax, Lanolin, Mineral Oil, Paraffin Wax & Phenoxyethanol.

Other Information

- do not use if inner seal is broken or missing

Learn more at www.HydrophorUS.com

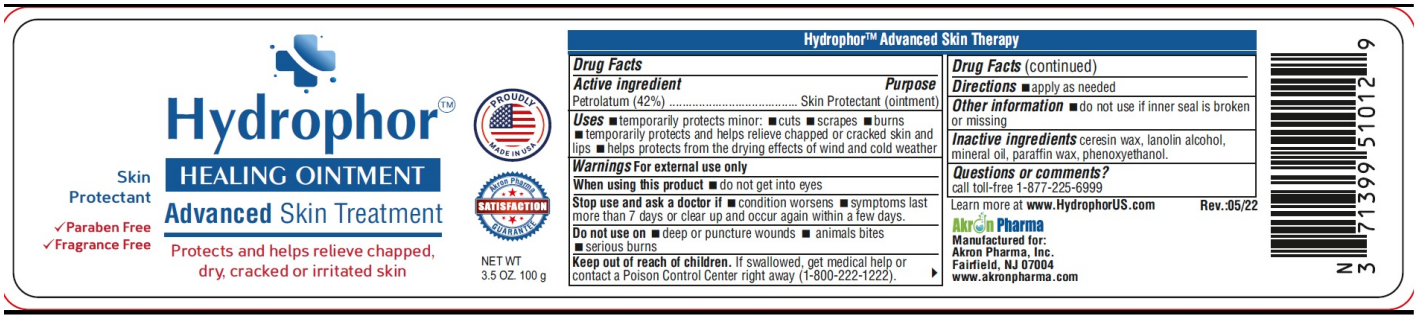
Questions?

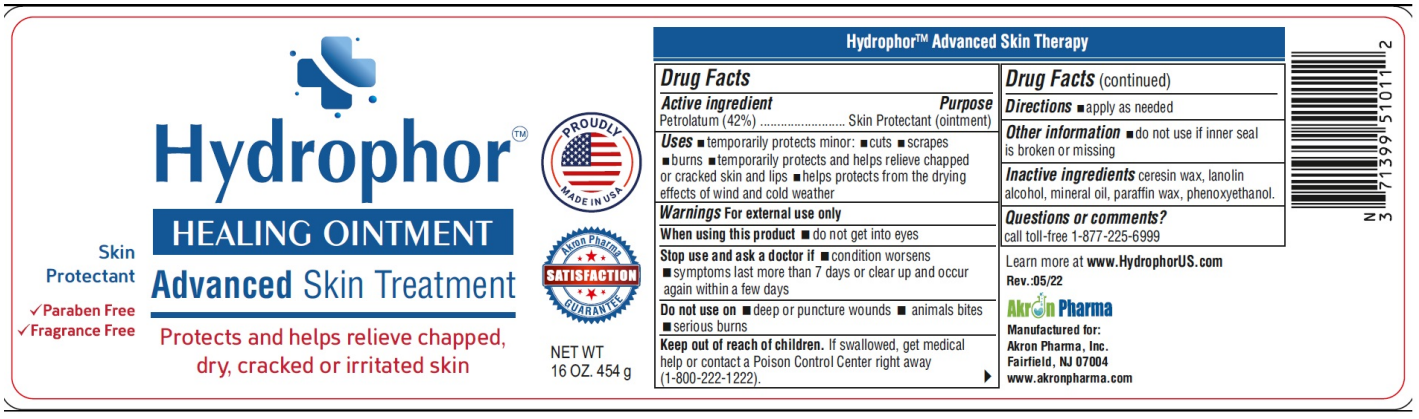
Please Call 1(877) 225-6999

Manufactured for:

Akron Pharma, Inc.
Fairfield, NJ 07004

Manufactured in U.S.A





HYDROPHOR

hydrophor ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71399-5101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	42 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CERESIN (UNII: Q1LS2UJO3A)	
LANOLIN (UNII: 7EV65EAW6H)	
MINERAL OIL (UNII: T5L8T28FGP)	
PARAFFIN (UNII: I9O0E3H2ZE)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-5101-1	454 g in 1 JAR; Type 0: Not a Combination Product	05/18/2021	
2	NDC:71399-5101-2	100 g in 1 JAR; Type 0: Not a Combination Product	05/18/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	05/18/2021	

Labeler - Akron Pharma Inc. (067878881)

Revised: 3/2023 Akron Pharma Inc.