

AMMENS MEDICATED POWDER ORIGINAL- zinc oxide powder

Idelle Labs, Ltd

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.

Ammens Medicated Powder

Medicated Powder

Ammens

Original Medicated Formula

All-Day Protection

- soothes
- deodorant protection
- relieves chafing
- cools

Active Ingredient Zinc Oxide 9.1%

Active Ingredient:

Zinc Oxide 9.1%

Purpose:

Skin protectant powder

Uses:

Dries the oozing and weeping of poison

- ivy
- oak
- sumac

Warning:

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

Keep out of reach of children.

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

Directions:

Apply as needed.

Other Information:

- Do not use on broken skin.
- When using this product, keep away from face and mouth to avoid breathing it in.

Inactive Ingredients:

Corn starch, fragrance, 8-hydroxyquinoline, 8-hydroxyquinoline sulfate, isostearic acid, PPG-20 methyl glucose ether, talc.

Questions or comments?

1-800-487-7273

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Made in the USA and distributed by Idelle Labs, Ltd., El Paso, TX 79912

Medicated
Powder

AMMENS[®]

**ORIGINAL
MEDICATED
FORMULA**

ALL-DAY PROTECTION



- Soothes
- Deodorant Protection
- Relieves Chafing
- Cools

AM 1058 H01 314

ACTIVE INGREDIENT • ZINC OXIDE 9.1%

NET WT. 11oz. (312g.)

label

AMMENS MEDICATED POWDER ORIGINAL

zinc oxide powder

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41595-1058
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	9.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
OXYQUINOLINE (UNII: 5UTX5635HP)	
ISOSTEARIC ACID (UNII: X33R8U0062)	
TALC (UNII: 7SEV7J4R1U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41595-1058-1	177 g in 1 CANISTER		
2	NDC:41595-1058-2	312 g in 1 CANISTER		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part347	02/01/2003	

Labeler - Idelle Labs, Ltd (128822926)

Registrant - Idelle Labs, Ltd (128822926)

Establishment

Name	Address	ID/FEI	Business Operations
PTI Union, LLC		807308858	MANUFACTURE(41595-1058)

Establishment

Name	Address	ID/FEI	Business Operations
M.K. Packaging, Inc.		047022405	MANUFACTURE(41595-1058)