

ALUMINUM ACETATE ASTRINGENT- calcium acetate and aluminum sulfate powder, for solution

TAGI Pharma Incorporated

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.

**ALUMINUM ACETATE
ASTRINGENT SOLUTION**

Drug Facts

Active ingredient (in each packet)

Aluminum acetate

(Each powder packet, when mixed in water and ready for use, provides the active ingredient aluminum acetate resulting from the reaction of calcium acetate 839 mg and aluminum sulfate 1191 mg.)

Purpose

Astringent

Uses

temporarily relieves minor skin irritations due to:

- poison ivy
- poison oak
- poison sumac
- insect bites
- athlete's foot
- rashes caused by soaps, detergents, cosmetics, or jewelry

Warnings

For external use only

When using this product

- avoid contact with eyes
- do not cover compress or wet dressing with plastic to prevent evaporation
- in some skin conditions, soaking too long may overdry

Stop use and ask a doctor if condition worsens or symptoms persist for more than 7 days

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

Directions

- mix one, two, or three packets in 16 oz of water to obtain the following modified Burrow's Solution

Number of Packets	Dilution	% Aluminum acetate
one packet	1:40 dilution	0.14%

two packets	1:20 dilution	0.28%
three packets	1:13 dilution	0.42%

- do not strain or filter the solution
- can be used as a compress, wet dressing, or a soak

FOR USE AS A COMPRESS OR WET DRESSING:

- soak a clean, soft cloth in the solution
- apply cloth loosely to affected area for 15 to 30 minutes
- repeat as needed or as directed by a doctor
- discard solution after each use

FOR USE AS A SOAK:

- soak affected area for 15 to 30 minutes as needed, or as directed by a doctor
- repeat 3 times a day or as directed by a doctor
- discard solution after each use

Other information

protect from excessive heat

Inactive ingredients

dextrin

Questions or comments?

1-888-EPIC-RX1

Distributed by:
TAGI Pharma
South Beloit, IL 61080

PRINCIPAL DISPLAY PANEL - 12 Powder Packets Carton

NDC 51224-153-24

**ALUMINUM ACETATE
ASTRINGENT SOLUTION**

Soothing, Effective Relief of Minor Skin Irritations due to:

- Poison Ivy**
- Athlete's Foot**
- Insect Bites**
- Rashes**

tagi PHARMA

Manufactured by:
Epic Pharma, LLC
Laurelton, NY 11413


Made in USA

Distributed by:
TAGI Pharma
South Beloit, IL 61080

12 POWDER PACKETS

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48041-0



Questions or comments? 1-800-EPIC-RX1

Inactive ingredients dextin

Other information protect from excessive heat

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- if some skin conditions, soaking too long may overly
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Uses temporarily relieves minor skin irritations due to:

- poison ivy ● poison oak ● insect bites ● athlete's foot
- rashes caused by soaps, detergents, cosmetics, or jewelry

Drug Facts

Active ingredient (in each packet) Aluminum acetate

Purpose Astringent

(Each powder packet, when mixed in water and ready for use, provides the active ingredient aluminum acetate resulting from the reaction of calcium acetate 859 mg and aluminum sulfate 1191 mg.)

LOT:

EXP:

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12 POWDER PACKETS

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NDC 51224-153-24

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Poison Ivy

Athlete's Foot

Insect Bites

Rashes

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ALUMINUM ACETATE ASTRINGENT
calcium acetate and aluminum sulfate powder, for solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51224-153
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Calcium Acetate (UNII: Y882YXF34X) (Calcium Cation - UNII:2M83C4R6ZB)	Calcium Acetate	839 mg in 2030 mg
Aluminum sulfate (UNII: 34S289N54E) (Aluminum Cation - UNII:3XHB1D032B)	Aluminum sulfate	1191 mg in 2030 mg

Inactive Ingredients

Ingredient Name	Strength
icodextrin (UNII: 2NX48Z0A9G)	

Product Characteristics

Color	WHITE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51224-153-24	12 in 1 CARTON		
1	NDC:51224-153-12	12 in 1 BOX		
1	NDC:51224-153-99	2030 mg in 1 POUCH		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part347	06/15/2011	

Labeler - TAGI Pharma Incorporated (963322560)

Revised: 6/2011

TAGI Pharma Incorporated