## ZO SKIN HEALTH ASTRINGENT SOLUTION- aluminum acetate powder ZO Skin Health, 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.

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# ZO® Skin Health Astringent Solution Powder National Drug Code 42851-090-06

#### **Drug Facts**

Active Ingredients	Purpose	
Aluminum Sulfate Tetradecahydrate, 2850 mg	Astringent*	
Calcium AcetateMonohydrate, 2016 mg	Astringent*	

<sup>\*</sup>When combined together in water, these ingredients form the active ingredient aluminum acetate. See **Directions** 

#### Uses

For temporary relief of minor skin irritations due to rashes caused by soaps, detergents, cosmetics, or jewelry.

## Warnings

## For external use only.

## When using this product

- avoid contact with eyes. If contact occurs, rinse thoroughly with water.
- 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**

## For powder dosage form:

- dissolve 1 packet in 12 oz. of cool or warm water
- stir until fully dissolved; do not strain or filter. The resulting mixture contains 0.45%

aluminum acetate and is ready for use.

#### 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

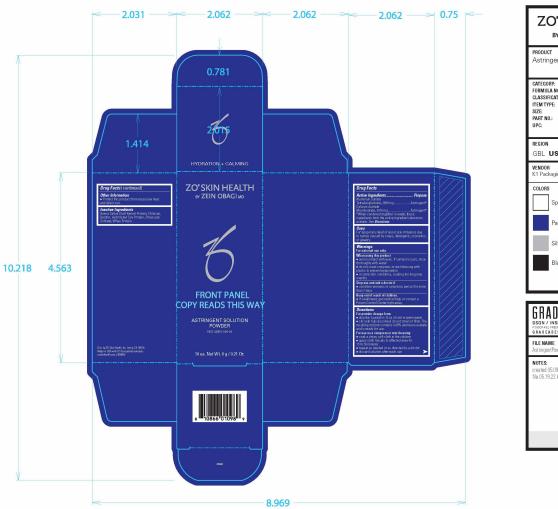
- repeat as needed or as directed by a doctor
- discard solution after each use

#### Other Information

■ Protect this product from excessive heat and direct sun

### Inactive Ingredients

Avena Sativa (Oat) Kernel Protein, Chitosan, Dextrin, Hydrolyzed Soy Protein, Potassium Sorbate, Whey Protein.





#### **ZO SKIN HEALTH ASTRINGENT SOLUTION**

aluminum acetate powder

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42851-090
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALUMINUM ACETATE (UNII: 80EHD8I43D) (ALUMINUM CATION - UNII: 3XHB1D032B)	ALUMINUM ACETATE	4866 mg in 6 g	

Inactive Ingredients		
Ingredient Name	Strength	
WHEY (UNII: 8617Z5FMF6)		
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)		
HYDROLYZED SOY PROTEIN (ENZYMATIC; 2000 MW) (UNII: 1394NXB9L6)		

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:42851-090- 06	6 g in 1 PACKET; Type 0: Not a Combination Product	02/21/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	02/21/2022	

## Labeler - ZO Skin Health, Inc. (826468527)

Revised: 10/2022 ZO Skin Health, Inc.