POLVO DE SULFA FIRST AID ANTIBIOTIC- bacitracin zinc, polymyxin b sulfate powder
GRANDALL DISTRIBUTING, LLC

Grandall (as PLD) - POLVO DE SULFA POWDER (48201-105)

Active Ingredients:

BACITRACIN ZINC 500 UNITS

POLYMYXIN B SULFATE 10,000 UNITS

Purpose:

First aid antibiotic

Uses

First aid to help prevent infection in minor
- cuts
- scrapes
- burns

Warnings
- For external use only

Do not use
- in the eyes
- if you are allergic to any of the ingredients
- over large areas of the body
- longer than 1 week unless directed by doctor

Ask a doctor before use if you have deep or puncture wounds, animal bites or serious burns

Stop use and ask a doctor if
- the conditions persists or gets worse
- a rash or other allergic reaction develops

Keep out of reach of childern.If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

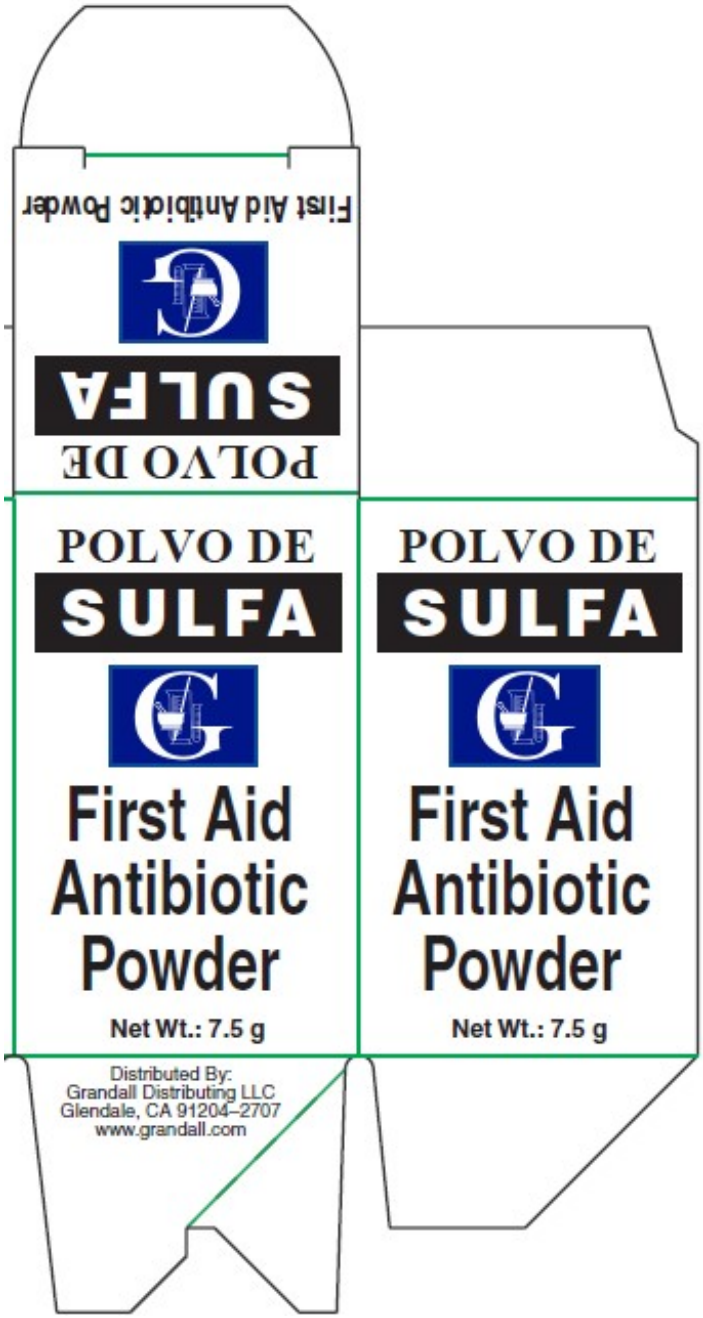
Adults and children 2 years of age and older:
- clean the affected area
- apply a light dusting of the powder on the area 1 to 3 times daily

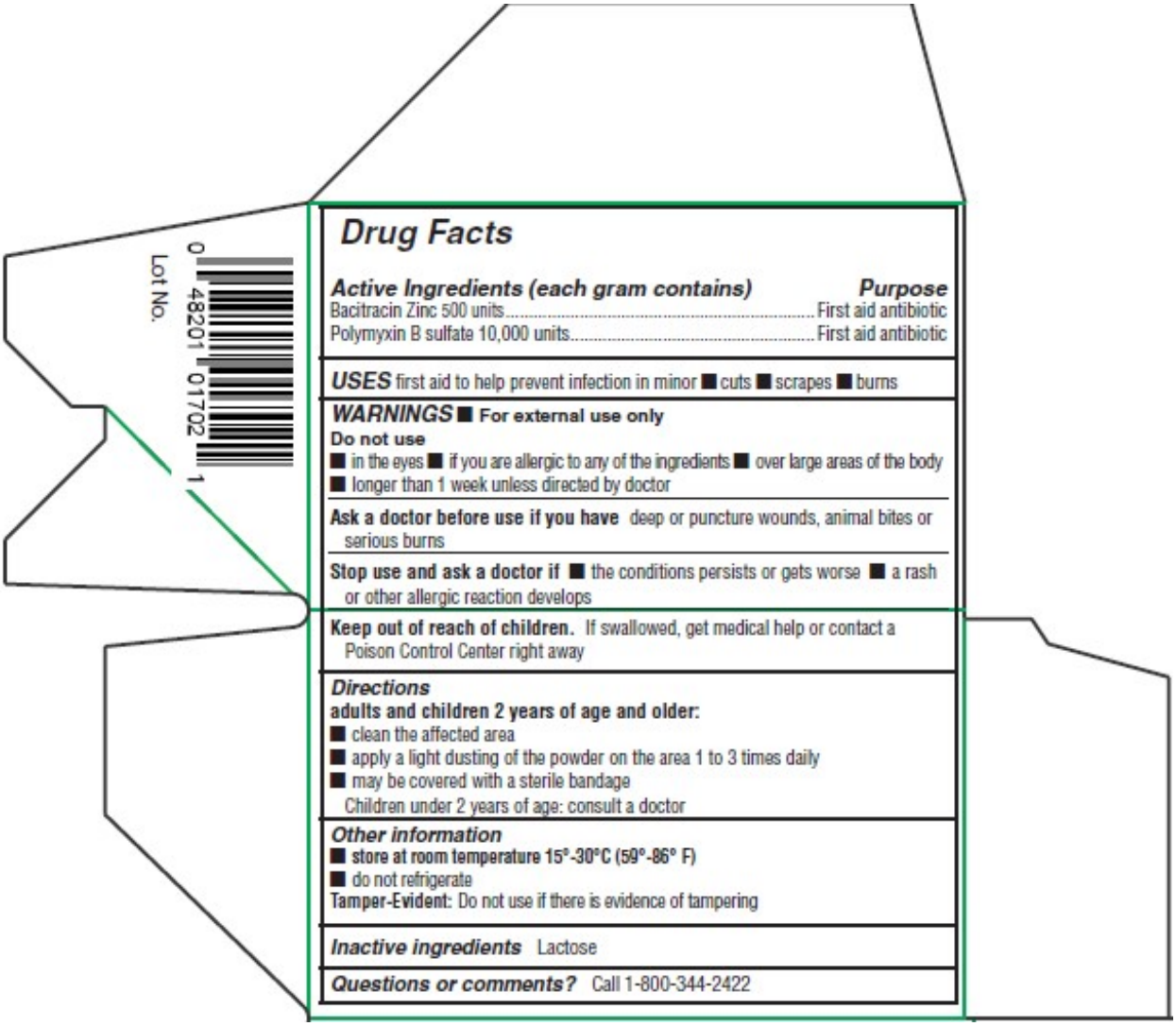
- may be covered with a sterile bandage

Children under 2 years of age: consult a doctor

Inactive Ingredient

Lactose





POLVO DE SULFA FIRST AID ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48201-105
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN ZINC	500 [USP'U] in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	10000 [USP'U] in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48201-105-75	1 in 1 BOX	07/18/2024	
1		7.5 g in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M004	07/18/2024	

Labeler - GRANDALL DISTRIBUTING, LLC (044428324)

Revised: 7/2024 GRANDALL DISTRIBUTING, LLC