TRIPLE ANTIBIOTIC- polymyxin b sulfate, neomycin sulfate, bacitracin, lidocaine hydrochloride ointment
Shandong Teking Pharmaceutical Co., 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.

Polymyxin B sulfate 5000units

Neomycin sulfate 3500units

Bacitracin 500units

Lidocaine hydrochloride 40mg

First aid antibiotic

First aid antibiotic

First aid antibiotic

External analgesic

First aid to help prevent infection and for the temporary relief of pain in minor:

- cuts
- scrapes
- burns
- surgical wound

For external use only

- If you are allergic to any of the ingredients
- In the eyes
- Over large areas of the body

- Deep or puncture wounds
- Animal bites
- Serious burns

- You need to use longer than 1 week
- Condition persists or gets worse
- Symptoms persist for more than 1 week, or clear up and occur again within a few days
- Rash or other allergic reaction develops

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

- Adults and children 2 years of age and older:

- Clean the affected area;

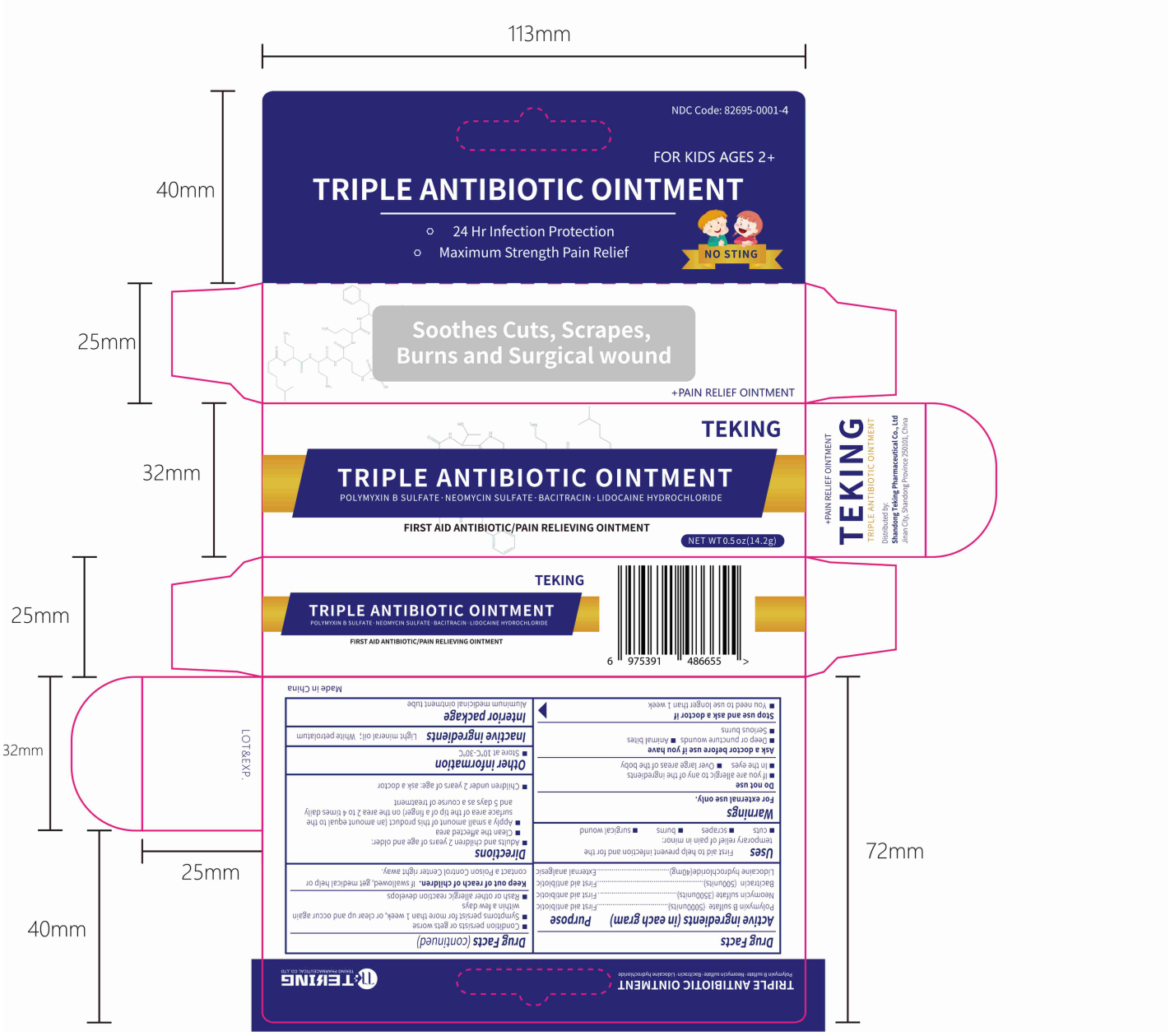
- Apply a small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 2 to 4 times daily and 5 days as a course of treatment.

- Children under 2 years of age: ask a doctor
- Store at 10℃-30℃

Light mineral oil

White petrolatum

TRIPLE ANTIBIOTIC OINTMENT



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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82695-0001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN (UNII: 58H6RWO52I) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [USP'U] in 1 g
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	40 mg in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	3500 [USP'U] in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [USP'U] in 1 g

Inactive Ingredients

Ingredient Name	Strength
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	

Product Characteristics

Color	white (White to light yellow suspension)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82695-0001-4	14.2 g in 1 TUBE; Type 0: Not a Combination Product	05/23/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	05/23/2022	

Labeler - Shandong Teking Pharmaceutical Co., Ltd (617652601)

Revised: 10/2022 Shandong Teking Pharmaceutical Co., Ltd