LEADER LIQUID BANDAGE- benzethonium chloride dyclonine hydrochloride liquid
LEADER/ Cardinal Health 110, Inc.

Leader Liquid Bandage

Active ingredients

Benzethonium chloride 0.2%
Dyclonine hydrochloride 0.75%

Purpose

First aid antiseptic
Topical analgesic

Uses

- first aid to prevent bacterial contamination in minor cuts, scrapes and burns
- for the temporary relief of pain

Warnings

For external use only.

Flammable:

Keep away from fire or flame.

Do not use

- in the eyes
- over large areas of the body
- longer than 1 week unless directed by a doctor

Ask a doctor before use if you have

- deep puncture wounds
- deep cuts
- animal bites
- serious burns

When using this product

- avoid contact with eyes

Stop use and ask a doctor if

- condition persists or gets worse
- symptoms persist for more than 7 days or clear up and occur again within a few days

- infection occurs

Keep out of reach of children.

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

Directions

- for adults and children 2 years of age and older
- clean affected area
- apply a small amount on the area 1-3 times daily and let dry
- a second coating may be applied for extra protection
- children under 2 years of age: consult a doctor

Other information

- **cap bottle tightly**
- store between 20° to 25°C (68° to 77°F)
- may temporarily sting upon application
- to remove, apply more Liquid Bandage and quickly wipe off

Inactive ingredients

acetone, amyl acetate, castor oil, ethyl acetate, nitrocellulose, SD alcohol 40

Questions

Call 1-866-964-0939

Principal Display Panel

LEADER

Liquid

Bandage

Benzethonium Chloride, 0.2%
First Aid Antiseptic

Dyclonine Hydrochloride, 0.75%
Topical Analgesic

Seals to Protect
For Small Cuts & Wounds

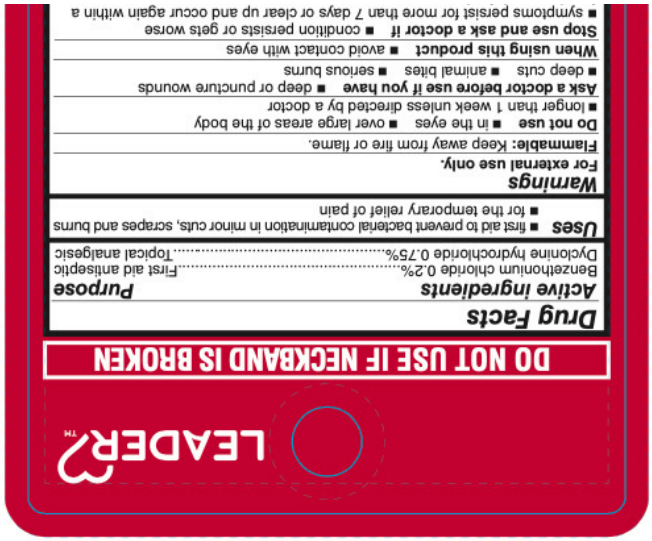
Prevents Infection

Invisible & Flexible

Water-Resistant

Built-In Applicator Brush

1.0 FL OZ (29.6 mL)



LEADER LIQUID BANDAGE

benzethonium chloride dyclonine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70000-0599
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.2 mg in 10 mL
DYCLONINE HYDROCHLORIDE (UNII: ZEC193879Q) (DYCLONINE - UNII:078A24Q30O)	DYCLONINE HYDROCHLORIDE	0.75 mg in 10 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ACETONE (UNII: 1364PS73AF)	
AMYL ACETATE (UNII: 92Q24NH7AS)	
CASTOR OIL (UNII: D5340Y2I9G)	
ETHYL ACETATE (UNII: 76845O8NMZ)	
PYROXYLIN (UNII: KYR8BR2X6O)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70000-0599-1	10 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	04/12/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	04/12/2022	

Labeler - LEADER/ Cardinal Health 110, Inc. (063997360)

Revised: 10/2026 LEADER/ Cardinal Health 110, Inc.