ORAL PAIN RELIEF GEL- benzocaine 20% gel Sled Distribution, LLC

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.

Oral Pain Relief Gel

Drug Facts

Active ingredient

Benzocaine 20% (w/w)

Purpose

Oral anesthetic

Uses

For the temporary relief of minor pain and sore mouth associated with toothache, minor dental procedures and irritations from dentures or orthodontic appliances.

Warnings

For oral use only.

stop use and ask a doctor if get any discomfort.

Stop use and ask a dentist or doctor if

- sore mouth does not improve in 7 days
- irritation, pain, or redness persists or worsens
- swelling, rash, fever or other allergic reaction develops

Keep out of reach of children.

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

Directions

Adults and children (2 years and older):

Dry affected area and apply medication undiluted. Use up to 4 times daily, but not more than every two hours, or as directed by a dentist or doctor.

Inactive ingredients

Polyethylene Glycol, Fragrance

Benzocaine 20% | Oral Anesthetic

NET WT 0.75 g

MY MEDIC[™]



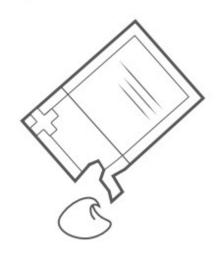


ORAL PAIN RELIEF GEL

Packet of oral pain relief gel to relieve tooth pain

ORAL PAIN RELIEF GEL Topical

Topical



EXPERIENCE ELEVATED FIRST AID

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NET WT .75 g - 1/38 oz



Drug Facts

Benzocaine 20% (w/w).....Oral Anesthetic

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Inactive Ingredients

Polyethylene Glycol, Fragrance.

Designed with with In Salt Lake City www.mymedic.com

MADE IN CHINA

Distributed By: SLED DISTRIBUTION LLC DBA MYMEDIC 120 North Redwood Rd. Ste. A, North Salt Lake , UT 84054





ORAL PAIN RELIEF GEL

benzocaine 20% gel

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:81417-121

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	200 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDWIA)	

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
NDC:81417-121-	1 in 1 PACKAGE	03/25/2022		
1	0.75 g in 1 BAG; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part356	03/25/2022	

Labeler - Sled Distribution, LLC (079772888)

Establishment				
Name	Address	ID/FEI	Business Operations	
Sled Distribution, LLC		079772888	manufacture(81417-121)	

Revised: 3/2022 Sled Distribution, LLC