# **TOPICALE XTRA-** benzocaine gel Medical Products Laboratories,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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#### **Topicale Xtra Gel - Jammin' Strawberry**

Benzocaine 20%

Oral Anesthetic

For the temporary relief of minor pain irritation associated with minor injury of the mouth and gums, canker sores, minor dental procedure, minor irritation caused by dentures or orthodontic appliances

#### Methemoglobinemia warning

Use of this product may cause methemoglobinemia, a serious condition that must be treated promptly because it reduces the amount of oxygen carried in blood. This can occur even if you have used this product before. Stop use and seek immediate medical attention if you or a child in your care develops pale, grey or blue colored skin (cyanosis), headache, rapid heart rate, shortness of breath, dizziness or light headedness, fatigue or lack of energy.

#### **Contraindications:**

Do not use in large quantities or over large areas of body Do not use for Teething Do not use in children under 2 years of age

## Allergy Alert:

Do not use if you have a history of allergy to local anesthetics such as benzocaine, butacaine, procaine or other "caine" anesthetics

# When using this product

Avoid contact with eyes

In case of accidental overdose, get medical help or contact a Poison Control Centre immediately.

Do not use more than directed.

## Stop use and ask a doctor

- 1. If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, swelling, rash, nausea or vomiting
- 2. If sore mouth symptoms do not improve in 7 days, or if irritation, pain or redness persists or worsens

Adults and children 12 years or older - Apply to the affected area. Allow to remain in place at least one minute and the spit out. Use up to 4 times daily or as directed by a

dentist or doctor.

Children 2-12 years of age - Should be supervised in the use of the product.

Children under 2 years of age - Do not use.

Do not use if imprinted seal under cap is broken or missing

Store at 68° to 77° F (20° - 25° C)

Benzalkonium Chloride (as a preservative), Carbomers, D&C Red Dye # 40, Flavorings, Polyethylene Glycol, Purified Water, Saccharin Sodium

Peel for Drug Facts

NDC 10733-176-01

Premier

Topicale Xtra

Topical Anesthetic Gel

Benzocaine, 20 %

REF 9007154 Jammin' Strawberry 28.35g (1 o.z.)

For Topical Use only - Not for Injection

Contains: 20 % Benzocaine in a specially designed glycol base

Made in U.S.A.

premierdentalco.com

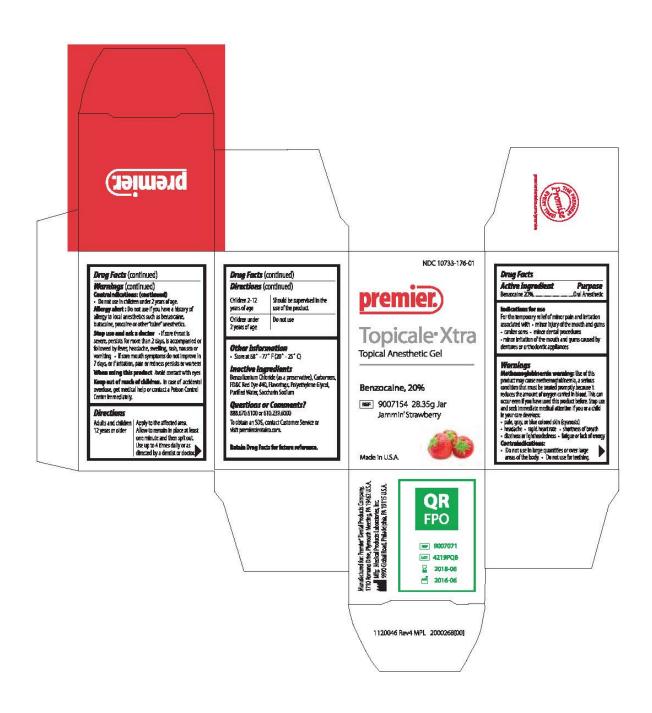
Manufactured for: Premier® Dental Products Company, 1710 Romano Drive, Plymouth Meeting, PA 19462 U.S.A.

Mfg: Medical Products Laboratories, Inc. 9990 Global Road Philadelphia, PA 19115 U.S.A.

1120049 MPL Rev5 . 2000267(00)

Questions or Comments?

888.670.6100 or 610.239.6000 M-Th: 7:30a.m. - 5:30p.m., F: 7:30a.m. - 4:00p.m. EST To obtain an SDS, contact Customer Service Department or visit premierdentalco.com.



Drug Facts (continued) **Directions** (continued)
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Children 2-12 years of age Should be supervised in the use of the product Children under 2 years of age Do not use

Other Information

Do not use if imprinted seal under cap is broken or missing
 Store at 68" - 77" F (20" - 25" C)

For Topical Use Only - Not for injection Contains: 20% Benzocalne in a specially designed glycol base.

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Peel for Drug Facts NDC 10733-176-01

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**Drug Facts** (continued)

Inactive Ingredients
Benallorium Chloride (as a preservative), Carbonners,
FD&C Red Dye #40, Randrings, Polyethylene Glycol, Purified
Water, Saccharin Sodium

**Questions or comments?** 888,670,5100 or 610,239,5000 M-Th: 7-30a.m. — 5:30p.m., P. 7-30 a.m. — 6:00p.m. EST

To obtain an SDS, contact Customer Service Department or visit premiers entail occur.

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Peel for Drug Facts NDC 10733-176-01 Topicale Xtra

Topical Anesthetic Gel Benzocaine, 20%

REF 9007154 Jammirf Strawberry 28.35g (1 oz.)

benzocaine gel

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:10733-176

Route of Administration ORAL

#### **Active Ingredient/Active Moiety**

FD&C RED NO. 40 (UNII: WZB9127XOA)

Ingredient Name	<b>Basis of Strength</b>	Strength
BENZOCAINE (UNII: U3RSY48IW5) (BENZOCAINE - UNII:U3RSY48IW5)	BENZ OCAINE	200 mg in 1 g

# Inactive Ingredients Ingredient Name Strength WATER (UNII: 059QF0KO0R) SACCHARIN SODIUM (UNII: SB8ZUX40TY) BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL OR ALLYL SUCROSE CROSSLINKED) (UNII: K6MOM3T5YL) POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	STRAWBERRY	Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:10733- 176-01	28.35 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/17/2022	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part356	08/17/2022		

# Labeler - Medical Products Laboratories, Inc. (002290302)