

**MAGIC PLUS ANESTHETIC- lidocaine hydrochloride, tetracaine hydrochloride,
and racepinephrine hydrochloride gel**
Tri-Lab Products Inc.

Magic Plus Topical Anesthetic Gel

ACTIVE INGREDIENTS

5.00% LIDOCAINE HYDROCHLORIDE, 2.00% TETRACAINE HYDROCHLORIDE, 0.02%
RACEPINEPHRINE HYDROCHLORIDE

INACTIVE INGREDIENTS

DEIONIZED WATER, HYDROXYETHYL CELLULOSE, SODIUM METABISULFITE, SODIUM
CHLORIDE, PROPYLENE GLYCOL, METHYLPARABEN, PROPYLPARABEN, DIAZOLIDINYL
UREA, AND TETRASODIUM EDTA

FOR EXTERNAL USE ONLY

For temporary relief of pain and swelling.

DIRECTIONS

Sensitivity test is strongly advised prior to use. Apply to area for temporary relief. Do not
use more than 2 times a day.

WARNINGS

CLIENT EYE PROTECTANT SUGGESTED

Do not get in the mouth or eyes. Do not use product if you are pregnant or a nursing
mother.

Keep out of the reach of children. If product comes into contact with the eyes wash
immediately. If accidentally swallowed seek immediate medical attention.

Do not use this product if you have any allergies to any of the product ingredients.

CAUTION

DISCONTINUE USE IF YOU HAVE SKIN IRRITATION OR SENSITIVITY.

Distributed by Trilab Products Inc.

PRINCIPAL DISPLAY PANEL - 28 G Bottle Label

Original!
Magic

Plus

TOPICAL ANESTHETIC GEL

Magic Works In Seconds!

RELIEVES PAIN
& SWELLING

1 FL OZ. (28 G)

FOR PROFESSIONAL USE ONLY

Store in a cool, dark place - Do not refrigerate!

Original! Magic Plus

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Distributed by Trilab Products Inc. 714.839.6543 NDC 84112-007-01

www.trilabproducts.com
Made in the USA Fountain Valley, CA 92708

MAGIC PLUS ANESTHETIC

lidocaine hydrochloride, tetracaine hydrochloride, and racepinephrine hydrochloride gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84112-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Lidocaine Hydrochloride (UNII: V13007Z41A) (Lidocaine - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	50 g in 1000 g
Tetracaine Hydrochloride (UNII: 5NF5D4OPCI) (Tetracaine - UNII:0619F35CGV)	Tetracaine Hydrochloride	20 g in 1000 g
Racepinephrine Hydrochloride (UNII: 336096P2WE) (Racepinephrine - UNII:GR0L9S3J0F)	Racepinephrine	200 mg in 1000 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
Sodium Chloride (UNII: 451W47IQ8X)	
Water (UNII: 059QF0KO0R)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Propylparaben (UNII: Z8IX2SC1OH)	
Methylparaben (UNII: A2I8C7HI9T)	
Diazolidinyl Urea (UNII: H5RIZ3MPW4)	
HYDROXYETHYL CELLULOSE (280 MPAS AT 2%) (UNII: 12VCE9HR9E)	
EDETATE SODIUM (UNII: MP1J8420LU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84112-007-01	1 in 1 BOX	04/01/2024	
1		28 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M017	04/01/2024	

Labeler - Tri-Lab Products Inc. (032851956)

Revised: 3/2024

Tri-Lab Products Inc.