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

Magic Plus Topical Anesthetic Spray

ACTIVE INGREDIENTS

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

INACTIVE INGREDIENTS

DEIONIZED WATER, 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. 714.839.6543.

PRINCIPAL DISPLAY PANEL - 28 G Bottle Label

Original!

Magic

Plus

TOPICAL ANESTHETIC SPRAY 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!



MAGIC PLUS ANESTHETIC

lidocaine hydrochloride, tetracaine hydrochloride, and racepinephrine hydrochloride liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84112-008	
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)	DACEDINIEDHDINIE	200 mg		

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: H5RIZ 3MPW4)		
EDETATE SODIUM (UNII: MP1J8420LU)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:84112- 008-01	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.