MAGIC PLUS ANESTHETIC- lidocaine hydrochloride, tetracaine hydrochloride, and racepinephrine hydrochloride gel Permanent Make Up Products LLC.

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. Store in a cool, dark place. Do not refrigerate.

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 immediatley. 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

Permanent Make Up Products LLC.

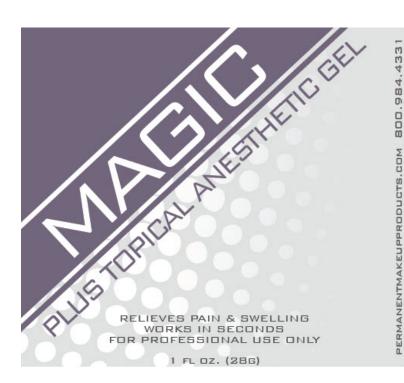
PRINCIPAL DISPLAY PANEL - 28 G Bottle Label

MAGIC

PLUS TOPICAL ANESTHETIC GEL RELIEVES PAIN & SWELLING WORKS IN SECONDS

FOR PROFESSIONAL USE ONLY

1 FL OZ. (28G)



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Distributed by Permanent Make Up Products LLC. Made in the USA Lake Havasu City, AZ 86403 NDC 84055-007-01



MAGIC PLUS ANESTHETIC

lidocaine hydrochloride, tetracaine hydrochloride, and racepinephrine hydrochloride gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84055-007
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 MPA.S AT 2%) (UNII: 12VCE9HR9E)	
EDETATE SODIUM (UNII: MP1J8420LU)	

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
NDC:84055- 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 - Permanent Make Up Products LLC. (030421491)

Revised: 3/2024 Permanent Make Up Products LLC.