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

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**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.

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

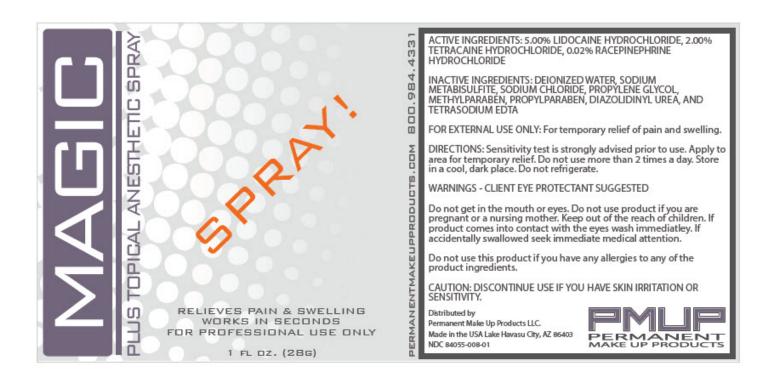
MAGIC
PLUS TOPICAL ANESTHETIC SPRAY

SPRAY!

RELIEVES PAIN & SWELLING WORKS IN SECONDS FOR PROFESSIONAL USE ONLY

1 FL OZ. (28G)

PERMANENTMAKEUPPRODUCTS.COM 800.984.4331



# **MAGIC PLUS ANESTHETIC**

lidocaine hydrochloride, tetracaine hydrochloride, and racepinephrine hydrochloride liquid

Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84055-008			
Route of Administration	TOPICAL					

Active Ingredient/Active Moiety				
Basis of Strength	Strength			
LIDOCAINE HYDROCHLORIDE ANHYDROUS	50 g in 1000 g			
Tetracaine Hydrochloride	20 g in 1000 g			
Racepinephrine	200 mg in 1000 g			
	LIDOCAINE HYDROCHLORIDE ANHYDROUS  Tetracaine Hydrochloride			

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)				

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

Revised: 3/2024 Permanent Make Up Products LLC.