NUMIT- benzocaine, lidocaine hydrochloride, and tetracaine hydrochloride liquid
Permanent Make Up Products LLC.

Numit Liquid

ACTIVE INGREDIENTS

5.00% BENZOCAINE HYDROCHLORIDE, 2.00% LIDOCAINE HYDROCHLORIDE, 2.00% TETRACAINE HYDROCHLORIDE

INACTIVE INGREDIENTS

ETHYL ALCOHOL, PROPYLENE GLYCOL, 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 - 14 G Bottle Label

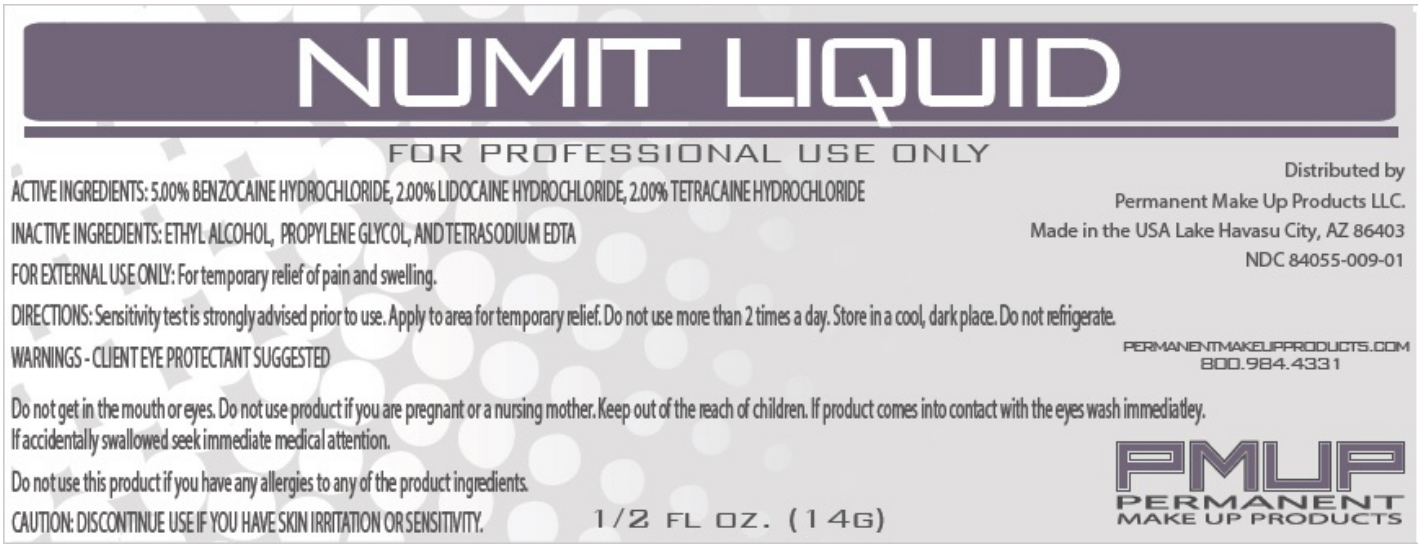
NUMIT LIQUID

FOR PROFESSIONAL USE ONLY

1/2 FL OZ. (14G)

PMUP

PERMANENT
MAKE UP PRODUCTS



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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84055-009
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Benzocaine (UNII: U3RSY48JW5) (Benzocaine - UNII:U3RSY48JW5)	Benzocaine	50 g in 1000 g
Lidocaine Hydrochloride (UNII: V13007Z41A) (Lidocaine - UNII:98PI200987)	Lidocaine Hydrochloride Anhydrous	20 g in 1000 g
Tetracaine Hydrochloride (UNII: 5NF5D4OPCI) (Tetracaine - UNII:0619F35CGV)	Tetracaine Hydrochloride	20 g in 1000 g

Inactive Ingredients	
Ingredient Name	**Strength**
Alcohol (UNII: 3K9958V90M)	500 g in 1000 g
Propylene Glycol (UNII: 6DC9Q167V3)	
Edetate Sodium (UNII: MP1J8420LU)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:84055-	1 in 1 BOX	04/01/2024	

1	009-01	1 in 1 BOX	04/01/2024	
1		14 g in 1 BOTTLE, GLASS; 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.