NUMQUICK ANALGESIC- epinephrine hydrochloride, lidocaine hydrochloride spray
Unit Dose, Ltd.

Numquick Analgesic

Drug Facts

Active ingredient

Epinephrine HCl 0.1 mg

Lidocaine HCl 50 mg

Purpose

Vasoconstrictor

Anesthetic

Uses:

Temporarily relieves local discomfort or pain and swelling or burning associated with anorectal disorders.

Warnings

External use only

certain persons can develop allergic reactions to the ingredients in this product **Allergy alert:**

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Do Not Use

this product in the rectum by using fingers or any mechanical device or applicator

When Using This Product

do not use more than directed.

Stop use and ask a doctor

if pain worsens or does not improve in 7 days
- redness, irritation, swelling, pain or other symptoms develop or increase
- if bleeding occurs

Keep out of the reach of children.

If accidentally swallowed get medical help or contact a Poison Control Center right away.

Directions:

Adults and Children over age 12 Spray 1-2ml to clean, dry affected area as needed up to 6 times daily. **---**

Children under age 12: ask a doctor

Store in a cool, dark place or refrigerate.

Inactive Ingredients:

Aqua (DI Water), Chlorobutanol, Glycerin, Sodium chloride, Sodium metabisulfite

Questions?

Call Toll Free 888/664-9990

Package Labeling:

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6/02

NUMQUICK ANALGESIC

epinephrine hydrochloride, lidocaine hydrochloride spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67194-008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

EPINEPHRINE HYDROCHLORIDE (UNII: WBB047OO38) (EPINEPHRINE - UNII:YKH834O4BH)	EPINEPHRINE HYDROCHLORIDE	0.1 mg in 1 mL
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	50 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CHLOROBUTANOL (UNII: HM4YQM8WRC)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67194-008-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/20/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	01/20/2016	

Labeler - Unit Dose, Ltd. (119080393)

Revised: 2/2024 Unit Dose, Ltd.