NUMQUICK TOPICAL ANALGESIC - epinephrine hydrochloride, lidocaine hydrochloride gel
Unit Dose, Ltd.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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Numquick Topical Analgesic

Drug Facts

Active ingredient
(-) Epinephrine HCL w/w 0.01%
Lidocaine HCl 5% w/w

Purpose
Vasoconstrictor
Local Anesthetic

Uses:
Temporarily relieves local discomfort or pain or burning associated with anorectal disorders.

Warnings:
External use only
Allergy alert: Certain persons can develop allergic reactions to the ingredients in this product.

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
• if redness, irritation, swelling, pain or other symptoms develop or increase
• if bleeding occurs

Keep out of reach of children.
If swallowed, get medical help or contact a Poison Control Center right away.

Directions:
• adults: clean the affected area with mild soap and warm water and rinse thoroughly
• dry gently
• apply 1-2 g. externally to the affected area up to 4 times daily
• children under 12 years: ask a doctor.
Other information:
Store in a cool dark place

Inactive ingredients:
Aqua (DI water), chlorobutanol, methylcellulose, sodium EDTA, sodium metabisulfite

Questions?
Call Toll Free 888-452-4946

Package Labeling:

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- Lidocaine HCl 5% w/w Local Anesthetic

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**NUMQUICK TOPICAL ANALGESIC**
epinephrine hydrochloride, lidocaine hydrochloride gel

**Product Information**

<table>
<thead>
<tr>
<th>Product Type</th>
<th>HUMAN OTC DRUG</th>
<th>Item Code (Source)</th>
<th>NDC:67194-005</th>
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<tbody>
<tr>
<td>Route of Administration</td>
<td>TOPICAL</td>
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**Active Ingredient/Active Moiety**

<table>
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<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
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<tbody>
<tr>
<td>EPINEPHRINE HYDROCHLORIDE (UNII: WBB047O038) (EPINEPHRINE - UNII:YKHB3404BH)</td>
<td>EPINEPHRINE</td>
<td>0.1 mg in 1 mL</td>
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**Inactive Ingredients**

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<thead>
<tr>
<th>Ingredient Name</th>
<th>Strength</th>
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<tbody>
<tr>
<td>WATER (UNII: 059QF0KO0R)</td>
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<tr>
<td>CHLOROBUTANOL (UNII: HM4YQM8WRC)</td>
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<tr>
<td>EDETATE SODIUM (UNII: MP1J8420LU)</td>
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<tr>
<td>SODIUM METABISULFITE (UNII: 4VON5FNS3C)</td>
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**Packaging**

<table>
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<tr>
<th>#</th>
<th>Item Code</th>
<th>Package Description</th>
<th>Marketing Start Date</th>
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<td>NDC:67194-005-01</td>
<td>59 mL in 1 BOTTLE; Type 0: Not a Combination Product</td>
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**Marketing Information**

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<tr>
<th>Marketing Category</th>
<th>Application Number or Monograph Citation</th>
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<tr>
<td>OTC monograph final</td>
<td>part346</td>
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**Labeler** - Unit Dose, Ltd. (119080393)

Revised: 12/2017