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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**Active ingredient**
Menthol 2%

**Purpose**
Topical Analgesic

**Uses**
For the temporary relief of minor aches and pains in muscles and joints associated with:

- simple backache
- sprains
- arthritis
- strains
- sports injuries
- bruises

**Warnings**
For external use only

**Do not use**
- with other topical pain relievers
- with heating pads or heating devices

**When using this product**
- do not use in or near eyes
- do not apply to wounds or damaged skin
- do not bandage tightly

**Stop use and ask a doctor if**
- condition worsens
- symptoms last for more than 7 days or clear up and occur again within a few days
- redness or irritation develops

**If pregnant or breast-feeding**
ask a health professional before use
Keep out of reach of children and pets. If swallowed, get medical help or contact a Poison Control Center right away.

Directions
- clean affected area before applying product
- adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily

Inactive Ingredients
- ammonium hydroxide, carboxymethylcellulose, cupric sulphate, FD&C blue no. 1, isopropyl alcohol, magnesium sulphate, sodium hydroxide, thymol, water

Package Label

REXALL ICE COLD ANALGESIC
menthol gel

Product Information

<table>
<thead>
<tr>
<th>Product Type</th>
<th>HUMAN OTC DRUG</th>
<th>Item Code (Source)</th>
<th>NDC:55910-088</th>
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<tbody>
<tr>
<td>Route of Administration</td>
<td>TOPICAL</td>
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Active Ingredient/Active Moiety

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<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
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<tbody>
<tr>
<td>MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)</td>
<td>MENTHOL</td>
<td>2 g in 100 g</td>
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Inactive Ingredients

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<thead>
<tr>
<th>Ingredient Name</th>
<th>Strength</th>
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<tr>
<td>CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)</td>
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<tr>
<td>CUPRIC SULFATE (UNII: LRX7AJ16DT)</td>
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<tr>
<td>MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)</td>
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<tr>
<td>THYMOL (UNII: 3J50XA376E)</td>
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<tr>
<td>FD&amp;C BLUE NO. 1 (UNII: H3R47K3TBD)</td>
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ISO PROPYL ALCOHOL (UNII: ND2M416302)
SODIUM HYDROXIDE (UNII: 55X04QC32I)
WATER (UNII: 059QF0KO0R)

Packaging

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<tr>
<th>#</th>
<th>Item Code</th>
<th>Package Description</th>
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<th>Marketing End Date</th>
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<td>227 g in 1 JAR; Type 0: Not a Combination Product</td>
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Marketing Information

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<th>Application Number or Monograph Citation</th>
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<td>OTC monograph not final</td>
<td>part348</td>
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Labeler - Dolgencorp, LLC (068331990)

Revised: 10/2018

Dolgencorp, LLC