MUCUS DM - guaifenesin and dextromethorphan hbr tablet, extended release
Amerisource Bergen

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Drug Facts

Active ingredients
(in each extended-release tablet)

Dextromethorphan Hydrobromide USP 60 mg
Guaifenesin USP 1200 mg

Purpose

Cough suppressant
Expectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily relieves:
  - cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
  - the intensity of coughing
  - the impulse to cough to help you get to sleep

Warnings

Do not use
- for children under 12 years of age
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson’s disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if you have

- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough accompanied by too much phlegm (mucus)

When using this product

- do not use more than directed

Stop use and ask a doctor if

- cough lasts more than 7 days, comes back, or occurs with fever, rash, or persistent headache. These
cough lasts more than 7 days, comes back, or occurs with fever, rash, or persistent headache. These could be signs of a serious illness.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- do not crush, chew, or break tablet
- take with a full glass of water
- this product can be administered without regard for timing of meals
- adults and children 12 years and older: 1 tablet every 12 hours; not more than 2 tablets in 24 hours
- children under 12 years of age: do not use

Other information

- tamper evident: do not use if carton is open or if printed seal on blister is broken or missing
- store at 20° to 25°C (68° to 77°F)

Inactive ingredients

colloidal silicon dioxide, hypromellose, magnesium stearate, microcrystalline cellulose, povidone, pregelatinized starch (maize)

Questions?

call 1-855-274-4122. You may also report side effects to this phone number.

*This product is not manufactured or distributed by Reckitt Benckiser, distributor of Maximum Strength Mucinex® DM.

Distributed By
AmerisourceBergen
1300 Morris Drive
Chesterbrook, PA 19087

Questions or Concerns?
www.mygnp.com

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 1200 mg/60 mg (14 Tablet Carton Label)

Compare to Maximum Strength Mucinex® DM
MUCUS DM
guaiifenesin and dextromethorphan hbr tablet, extended release
Product Information

Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:46122-420
Route of Administration | ORAL

Active Ingredient/Active Moiety

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
</tr>
</thead>
<tbody>
<tr>
<td>GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)</td>
<td>GUAIFENESIN</td>
<td>1200 mg</td>
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<tr>
<td>DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)</td>
<td>DEXTROMETHORPHAN HYDROBROMIDE</td>
<td>60 mg</td>
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Inactive Ingredients

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Strength</th>
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<tbody>
<tr>
<td>SILICON DIOXIDE (UNII: ETJ7Z6XBU4)</td>
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<tr>
<td>HYPRO MELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)</td>
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<tr>
<td>MAGNESIUM STEARATE (UNII: 70097M630)</td>
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<td>MICROCRYSTALLINE CELLULOSE (UNII: OPIR32D61U)</td>
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<td>Povidone K90 (UNII: RDB86HJVSZ)</td>
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<tr>
<td>Povidone K25 (UNII: K0KQV10C35)</td>
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<tr>
<td>STARCH, CORN (UNII: O8232NY3SJ)</td>
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Product Characteristics

| Color | WHITE (White to Off-white) |
| Shape | OVAL |
| Flavor | |
| Contains | |

Score no score
Size 22mm
Imprint Code X;63

Packaging

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<th>#</th>
<th>Item Code</th>
<th>Package Description</th>
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<th>Marketing End Date</th>
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<tbody>
<tr>
<td>1</td>
<td>NDC:46122-420-74</td>
<td>2 in 1 CARTON</td>
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<tr>
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<td>7 in 1 BLISTER PACK; Type 0: Not a Combination Product</td>
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<tr>
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Marketing Information

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<td>03/17/2017</td>
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Labeler - Amerisource Bergen (007914906)

Registrant - Aurohealth LLC (078728447)
<table>
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<tr>
<th>Name</th>
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<th>ID/FEI</th>
<th>Business Operations</th>
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<tr>
<td>Aurobindo Pharma Limited</td>
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<td>650381903</td>
<td>ANALYSIS(46122-420), MANUFACTURE(46122-420)</td>
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