MUCINEX DM- guaifenes in and dextromethorphan hydrobromide tablet, extended release RB Health (US) LLC

Mucinex®DM

Drug Facts

| Active ingredients (in each extended-release bilayer tablet) | Purposes |
|--|-------------------|
| Dextromethorphan HBr 30 mg | Cough suppressant |
| Guaifenesin 600 mg | 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 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 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 or 2 tablets every 12 hours; not more than 4 tablets in 24 hours
- children under 12 years of age: do not use

Other information

■ store at 20-25°C (68-77°F)

Inactive ingredients

carbomer homopolymer type B; D&C yellow #10 aluminum lake; hypromellose, USP; magnesium stearate, NF; microcrystalline cellulose, NF; sodium starch glycolate, NF

Questions?

1-866-MUCINEX (1-866-682-4639)

You may also report side effects to this phone number.

Dist. by: RB Health (US) Parsippany, NJ 07054-0224

Made in England

PRINCIPAL DISPLAY PANEL - 20 Tablet Blister Pack Carton

NDC 63824-056-32

Mucinex®DM

600 mg guaifenesin & 30 mg dextromethorphan HBr extended-release bi-layer tablets

EXPECTORANT & COUGH SUPPRESSANT

12 HOUR_®

- Controls Cough
- Thins and Loosens Mucus
- ☐ Immediate and Extended Release

20 EXTENDED-RELEASE BI-LAYER TABLETS MucinexDM

Tamper evident: Do not use if carton is open or if printed seal on blister is broken or missing.

EXTENDED-RELEASE BI-LAYER TABLETS

NDC 63824-056-32

Mucinexida

600 mg guaifenesin & 30 mg dextromethorphan HBr extended-release bi-layer tablets

EXPECTORANT & COUGH SUPPRESSANT



- ✓ Controls Cough
- ✓ Thins and Loosens Mucus
- ✓ Immediate and Extended Release
- 20 EXTENDED-RELEASE BI-LAYER TABLETS

Choose from the MUCINEX® line of products to treat your sympton

www.StopMedicineAbuse.org

63824 05620

3077437

Pharma Code

Glue Line



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EXPECTORANT & COUGH SUPPRESSANT

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Drug Facts (continued)

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Dist. by: RB Health (US)

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Patents: www.rb.com/patents 011119 3077437

EXPECTOBANT & COUGH SUPPRESSANT

mmediate-Release

Extended-Rélease



MUCINEX DM

| Product Information | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:63824-056 |
| Route of Administration | ORAL | | |

| Active Ingredient/Active Moiety | | | |
|--|----------------------------------|----------|--|
| Ingredient Name | Basis of Strength | Strength | |
| Guaifenesin (UNII: 495W7451VQ) (Guaifenesin - UNII:495W7451VQ) | Guaifenesin | 600 mg | |
| Dextromethorphan Hydrobromide (UNII: 9D2RTI9KYH) (Dextromethorphan - UNII:7355X3ROTS) | Dextromethorphan Hydrobromide | 30 mg | |

| Inactive Ingredients | | | | |
|--|----------|--|--|--|
| Ingredient Name | Strength | | | |
| Carbomer Homopolymer Type B (Allyl Pentaerythritol Crosslinked) (UNII: HHT01ZNK31) | | | | |
| D&C yellow no. 10 (UNII: 35SW5USQ3G) | | | | |
| aluminum oxide (UNII: LMI26O6933) | | | | |
| HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29 V3WO) | | | | |
| magnesium stearate (UNII: 70097M6I30) | | | | |
| MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U) | | | | |

| Product Characteristics | | | |
|-------------------------|--------------------------|--------------|-------------|
| Color | WHITE (yellow and white) | Score | no score |
| Shape | OVAL | Size | 16 mm |
| Flavor | | Imprint Code | Mucinex;600 |
| Contains | | | |

| Packaging | | | | |
|-----------|----------------------|---|-----------------------------|---------------------------|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:63824-056- 89 | 4 in 1 CARTON | 03/30/2018 | |
| 1 | | 17 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |
| 2 | NDC:63824-056-36 | 1 in 1 CARTON | 06/26/2012 | |
| 2 | | 6 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |
| 3 | NDC:63824-056-32 | 1 in 1 CARTON | 06/26/2012 | |
| 3 | | 20 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |
| 4 | NDC:63824-056-34 | 2 in 1 CARTON | 06/26/2012 | |
| 4 | | 20 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |
| 5 | NDC:63824-056- 69 | 3 in 1 CARTON | 06/26/2012 | |
| 5 | | 20 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |
| 6 | NDC:63824-056-74 | 24 in 1 CARTON | 06/26/2012 | |
| 6 | NDC:63824-056-73 | 2 in 1 POUCH; Type 0: Not a Combination Product | | |
| 7 | NDC:63824-056-72 | 2 in 1 POUCH; Type 0: Not a Combination Product | 06/26/2012 | |

| 8 | NDC:63824-056-11 | 2 in 1 CARTON | 06/26/2012 | |
|---|------------------|---|------------|--|
| 8 | | 12 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |
| 9 | NDC:63824-056-01 | 5 in 1 CARTON | 06/26/2012 | |
| 9 | | 20 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |

| Marketing Information | | | |
|-----------------------|--|----------------------|--------------------|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| NDA | NDA021620 | 06/26/2012 | |
| | | | |

Labeler - RB Health (US) LLC (081049410)

Revised: 10/2019 RB Health (US) LLC