

MEIJER MUCUS RELIEF DM MAXIMUM STRENGTH- dextromethorphan hbr and guaifenesin solution
MEIJER, INC.

Meijer Mucus Relief DM Maximum Strength

Drug Facts

| <i>Active ingredients (in each 20 mL)</i> | <i>Purposes</i> |
|--|------------------------|
| Dextromethorphan HBr 20 mg | Cough suppressant |
| Guaifenesin 400 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

- 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 that occurs with 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 that lasts. These could be signs of a serious condition.

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 at 1-800-222-2222.

Directions

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided
- do not use dosing cup with other products
- dose as follows or as directed by a doctor
- mL = milliliter
- **adults and children 12 years and older:** 20 mL every 4 hours
- **children under 12 years of age:** Do not use

Other information

- **each 20 mL contains:** sodium 8 mg
- low sodium
- store at room temperature
- do not refrigerate
- dosing cup provided

Inactive ingredients

anhydrous citric acid, edetate disodium, FD&C Blue No. 1, FD&C Red No. 40, flavors, potassium citrate, propylene glycol, propyl gallate, purified water, sodium benzoate, sorbitol, sucralose, xanthan gum.

Questions or comments?

1-866-467-2748

PRINCIPAL DISPLAY PANEL

NDC# 41250-748-06

Compare to Maximum Strength Mucinex[®] Fast-Max[®] DM max active ingredients

mucus relief DM

Dextromethorphan HBr/Cough Suppressant
Guaifenesin/Expectorant

Maximum Strength

- **Controls Cough**

- Relieves Chest Congestion
- Thins & Loosens Mucus
- 4 Hour Dosing

For Ages 12+

6 FL OZ (180 mL)

Tamper evident: Do not use if printed inner seal under cap is broken or missing.

DIST. BY MEIJER DISTRIBUTION INC.

GRAND RAPIDS, MI 49544

www.meijer.com

*This product is not manufactured or distributed by Reckitt BRB Health (US) LLC, owner of the registered trademark Mucinex® and Fast -Max®



MEIJER MUCUS RELIEF DM MAXIMUM STRENGTH

dextromethorphan hbr and guaifenesin solution

Product Information

| | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:41250-748 |
| Route of Administration | ORAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|-----------------|-------------------|----------|
|-----------------|-------------------|----------|

| | | |
|--|-------------------------------|--------------------|
| dextromethorphan hydrobromide (UNII: 9D2RTI9KYH) (dextromethorphan - UNII:7355X3ROTS) | dextromethorphan hydrobromide | 20 mg in 20 mL |
| guaifenesin (UNII: 495W7451VQ) (guaifenesin - UNII:495W7451VQ) | guaifenesin | 400 mg in 20 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| anhydrous citric acid (UNII: XF417D3PSL) | |
| edetate disodium (UNII: 7FLD91C86K) | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |
| FD&C red No. 40 (UNII: WZB9127XOA) | |
| POTASSIUM CITRATE (UNII: EE90ONI6FF) | |
| propylene glycol (UNII: 6DC9Q167V3) | |
| propyl gallate (UNII: 8D4SNN7V92) | |
| water (UNII: 059QF0KO0R) | |
| sodium benzoate (UNII: OJ245FE5EU) | |
| sorbitol (UNII: 506T60A25R) | |
| sucralose (UNII: 96K6UQ3ZD4) | |
| xanthan gum (UNII: TTV12P4NEE) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:41250-748-06 | 180 mL in 1 BOTTLE; Type 0: Not a Combination Product | 04/20/2020 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M012 | 04/20/2020 | |

Labeler - MEIJER, INC. (006959555)

Revised: 11/2023

MEIJER, INC.