MUCUS RELIEF- guaifenesin 400 mg tablet NewVue LLC

Mucus Relief Guaifenesin 400 mg Caplets

Active ingredient (in each tablet)

Guaifenesin 400 mg

Purpose

Expectorant

Uses

 helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive to rid the bronchial passageway of bothersome mucus

Warnings

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 excessive phlegm (mucus)

When using this product

- do not exceed recommended dosage
- do not use for more than 7 days

Stop use and ask a doctor if

 cough lasts for more than 7 days, recurs, or is accompanied by fever, rash, or persistent headache.

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.

Directions:

- adults and children 12 years of age and over:take 1 tablet every 4 hours with a full glass of water while symptoms persist. Do not exceed 6 doses in 24 hours.
- children under 12 years:do not use

Other information

- each tablet contains: sodium 1.24 mg VERY LOW SODIUM
- store at 25°C (77°F) excursions between 15°-30°C (59°-86°F)
- keep in a dry place and do not expose to heat
- read all product information before using
- TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Inactive ingredients

colloidal silicon dioxide, magnesium stearate, microcrystalline cellulose, stearic acid, sodium starch glycolate

Questions or Comments

www.getnewvue.com

Principal Display Panel

300 ct



MUCUS RELIEF

guaifenesin 400 mg tablet

| Product Information | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:83813-003 |
| Route of Administration | ORAL | | |

| Active Ingredient/Active Moiety | | |
|--|-------------------|----------|
| Ingredient Name | Basis of Strength | Strength |
| GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) | GUAIFENESIN | 400 mg |

| Inactive Ingredients | | |
|--|----------|--|
| Ingredient Name | Strength | |
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4) | | |
| MAGNESIUM STEARATE (UNII: 70097M6I30) | | |
| CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U) | | |
| STEARIC ACID (UNII: 4ELV7Z65AP) | | |
| POVIDONE (UNII: FZ989GH94E) | | |
| MALTODEXTRIN (UNII: 7CVR7L4A2D) | | |
| CROSCARMELLOSE SODIUM (UNII: M280L1HH48) | | |

| Product Characteristics | | | |
|-------------------------|-------|--------------|----------|
| Color | white | Score | no score |
| Shape | OVAL | Size | 17mm |
| Flavor | | Imprint Code | S400 |
| Contains | | | |

| ı | Packaging | | | | |
|---|-----------|----------------------|--|-------------------------|-----------------------|
| | # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| | 1 | NDC:83813-003- 03 | 300 in 1 BOTTLE; Type 0: Not a Combination Product | 07/01/2020 | |

| Marketing Information | | | |
|-----------------------|---|-------------------------|-----------------------|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC Monograph Drug | M012 | 04/13/2016 | |
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Labeler - NewVue LLC (119120572)

Revised: 4/2024 NewVue LLC