DAYTIME MUCUS RELIEF SEVERE COLD AND NIGHTTIME COLD AND FLU MAXIMUM STRENGTH- acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl,guaifenesin CVS Pharmacy

Drug Facts

Active ingredients in Daytime (in each softgel) Acetaminophen 325 mg

Dextromethorphan HBr 10 mg
Guaifenesin 200 mg
Phenylephrine HCl 5 mg

Active ingredients in Nighttime (in each softgel) Acetaminophen 325 mg

Dextromethorphan HBr 10 mg
Doxylamine succinate 6.25 mg
Phenylephrine HCI 5 mg

Purpose for Daytime

Pain reliever/fever reducer

Cough suppressant

Expectorant

Nasal decongestant

Purpose for Nighttime

Pain reliever/fever reducer

Cough suppressant

Antihistamine

Nasal decongestant

Uses

DAYTIME

- temporarily relieves these common cold and flu symptoms
 - headache

- nasal congestion
- sore throat
- cough
- minor aches and pains
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily reduces fever

NIGHTTIME

- temporarily relieves these common cold and flu symptoms
 - cough
 - headache
 - minor aches and pains
 - sore throat
 - nasal congestion
 - runny nose and sneezing
 - controls cough to help you get to sleep
 - temporarily reduces fever

Warnings

DAYTIME and NIGHTTIME

Liver warning: These products contain acetaminophen. Severe liver damage may occur if you take:

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away

Sore throat warning: If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

Do not use

DAYTIME and NIGHTTIME

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- 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

DAYTIME

- liver disease
- diabetes
- high blood pressure
- heart disease
- thyroid disease
- trouble urinating due to an enlarged prostate gland
- a persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough that occurs with too much phlegm (mucus)

NIGHTTIME

- liver disease
- diabetes
- high blood pressure
- heart disease
- glaucoma
- thyroid disease
- trouble urinating due to an enlarged prostate gland
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough that occurs with too much phlegm (mucus

Ask a doctor or pharmacist before use if you are

DAYTIME

taking the blood thinning drug warfarin

NIGHTTIME

taking the blood thinning drug warfarin taking sedatives or tranquilizers

When using this product,

DAYTIME

do not use more than directed

NIGHTTIME

- do not use more than directed
- excitability may occur, especially in children
- marked drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

DAYTIME and NIGHTTIME

- nervousness, dizziness, or sleeplessness occur
- pain, nasal congestion, or cough gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back or occurs with rash or headache that lasts. These could be signs
 of a serious condition.

If pregnant or breast-feeding,

DAYTIME and NIGHTTIME

ask a health professional before use.

Keep out of reach of children.

DAYTIME and NIGHTTIME

Overdose warning: Taking more than the recommended dose can cause liver damage. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

DAYTIME

- do not take more than directed (see Overdose warning)
- do not take more than 12 softgels (Daytime and NightTime) in any 24-hour period
- adults and children 12 years of age and older: take 2 softgels every 4 hours
- children under 12 years of age: do not use
- when using other Daytime or Nighttime products, carefully read each label to ensure correct dosing

NIGHTTIME

- do not take more than directed (see Overdose warning)
- do not take more than 12 softgels (Daytime and Nighttime) in any 24-hour period
- adults and children 12 years of age and older: take 2 softgels every 4 hours
- children under 12 years of age: do not use
- when using other Daytime or Nighttime products, carefully read each label to ensure correct dosing

Other information

DAYTIME and **NIGHTTIME**

- store between 20-25°C (68-77°F)
- avoid excessive heat
- swallow whole, do not crush, chew, or dissolve

Inactive ingredients

DAYTIME

FD&C red #40, FD&C yellow #6, gelatin, glycerin, mannitol, polyethylene glycol, povidone, propylene glycol, purified water, sorbitan, sorbitol, titanium dioxide

NIGHTTIME

D&C yellow #10, FD&C blue #1, gelatin, glycerin, mannitol, polyethylene glycol, povidone, propylene glycol, purified water, sorbitan, sorbitol, titanium dioxide

Questions or comments?

DAYTIME and NIGHTTIME

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal Display Panel

DAY & NIGHT COMBO PACK

Compare to the active ingredients in Maximum Strength Mucinex® FAST-Max® Day Severe Cold Liquid Gels and Night Cold & Flu Liquid Gels†

DAYTIME

MAXIMUM STRENGTH

Daytime Severe Cold

Acetaminophen 325 mg Pain Reliever; Fever Reducer

Dextromethorphan HBr 10 mg Cough Suppressant

Guaifenesin 200 mg Expectorant

Phenylephrine HCL - 5 mg Nasal Decongestant

MULTI-SYMPTOM

- Relieves cough
- Thins & loosens mucus
- Relieves nasal & chest congestion
- Relieves aches, fever & sore throat

FOR AGES 12+

SOFTGELS**

(Liqui-filled capsules)

NIGHTTIME

MAXIMUM STRENGTH

Nighttime Cold & Flu

Acetaminophen 325 mg Pain reliever, Fever reducer

Dextromethorphan HBr 10 mg Cough suppressant

Doxylamine Succinate 6.25 mg Antihistamine

Phenylephrine HCL 5 mg Nasal decongestant

MULTI-SYMPTOM

- Relieves cough
- Relieves runny nose & sneezing
- Relieves nasal congestion
- Relieves aches, fever & sore throat

FOR AGES 12+

SOFTGELS**

(*Liqui-Filled Capsules)

*This product is not manufactured or distributed by Reckitt Benckiser, distributor of Maximum Strength Mucinex® Fast-Max® Day Severe Cold and Maximum Strength Mucinex® Fast-Max® Night Cold & Flu Liquid Gels.

TAMPER EVIDENT: DO NOT USE IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Distributed by: CVS Pharmacy, Inc.

One CVS Drive, Woonsocket, RI 02895

Product Label



CVS HEALTH Maximum Strength Daytime Severe Cold, Maximum Strength Nighttime Severe Cold

DAYTIME MUCUS RELIEF SEVERE COLD AND NIGHTTIME COLD AND FLU MAXIMUM STRENGTH

acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl, guaifenesin kit

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:69842-906

| F | Packaging | | | | |
|---|----------------------|---|-------------------------|-----------------------|--|
| # | t Item Code | Package Description | Marketing Start Date | Marketing End Date | |
| 1 | NDC:69842-906- 24 | 1 in 1 KIT; Type 0: Not a Combination Product | 11/29/2019 | 11/29/2026 | |

| Quantity of Parts | | | | |
|-------------------|------------------|------------------------|--|--|
| Part # | Package Quantity | Total Product Quantity | | |
| Part 1 | 8 BLISTER PACK | 8 | | |
| Part 2 | 16 BLISTER PACK | 16 | | |
| | | | | |

Part 1 of 2

NIGHTTIME COLD AND FLU MAXIMUM STRENGTH

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hcl capsule

Product Information

Route of Administration ORAL

| Active Ingredient/Active Moiety | | | | |
|--|----------------------------------|----------|--|--|
| Ingredient Name | Basis of Strength | Strength | | |
| ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) | ACETAMINOPHEN | 325 mg | | |
| DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 10 mg | | |
| DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL) | DOXYLAMINE SUCCINATE | 6.25 mg | | |
| PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS 297W6MV) | PHENYLEPHRINE HYDROCHLORIDE | 5 mg | | |

| Inactive Ingredients | | | |
|---|----------|--|--|
| Ingredient Name | Strength | | |
| D&C YELLOW NO. 10 (UNII: 35SW5USQ3G) | | | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | | | |
| GELATIN (UNII: 2G86QN327L) | | | |
| GLYCERIN (UNII: PDC6A3C0OX) | | | |
| POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) | | | |
| POVIDONE (UNII: FZ989GH94E) | | | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | | | |
| WATER (UNII: 059QF0KO0R) | | | |
| SORBITAN (UNII: 6092ICV9RU) | | | |
| SORBITOL (UNII: 506T60A25R) | | | |
| MANNITOL (UNII: 3OWL53L36A) | | | |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP) | | | |

| Product Characteristics | | | | |
|-------------------------|---------|--------------|----------|--|
| Color | green | Score | no score | |
| Shape | CAPSULE | Size | 20mm | |
| Flavor | | Imprint Code | 42A | |
| Contains | | | | |

| Pa | Packaging | | | | |
|----------------------------|-----------|--|-----------------------|--|--|
| # Item Package Description | | Marketing Start Date | Marketing End Date | | |
| 1 | | 8 in 1 CARTON | | | |
| | | 1 in 1 BLISTER PACK; Type 0: Not a Combination | | | |

| Marketing | Application Number or Monograph | Marketing Start | Marketing End |
|--------------------|---------------------------------|-----------------|---------------|
| Category | Citation | Date | Date |
| OTC Monograph Drug | M012 | 11/29/2019 | |

Part 2 of 2

DAYTIME SEVERE COLD MAXIMUM STRENGTH

acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl capsule

Product Information

Route of Administration ORAL

| Active Ingredient/Active Moiety | | | | |
|--|----------------------------------|----------|--|--|
| Ingredient Name | Basis of Strength | Strength | | |
| ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) | ACETAMINOPHEN | 325 mg | | |
| DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 10 mg | | |
| GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) | GUAIFENESIN | 200 mg | | |
| PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV) | PHENYLEPHRINE HYDROCHLORIDE | 5 mg | | |

| Inactive Ingredients | |
|---|----------|
| Ingredient Name | Strength |
| FD&C RED NO. 40 (UNII: WZB9127XOA) | |
| FD&C YELLOW NO. 6 (UNII: H77VEI93A8) | |
| GELATIN (UNII: 2G86QN327L) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) | |
| POVIDONE (UNII: FZ989GH94E) | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | |
| WATER (UNII: 059QF0KO0R) | |
| SORBITAN (UNII: 6092ICV9RU) | |
| SORBITOL (UNII: 506T60A25R) | |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP) | |
| MANNITOL (UNII: 3OWL53L36A) | |

| Product Characteristics | | | | |
|-------------------------|---------|-------|----------|--|
| Color | orange | Score | no score | |
| Shape | CAPSULE | Size | 20mm | |

| Flavor | Impr | int Code | 12A |
|----------|------|----------|-----|
| Contains | | | |

| Packaging | | | | | | | |
|-----------|--------------|--|-------------------------|-----------------------|--|--|--|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date | | | |
| 1 | | 16 in 1 CARTON | | | | | |
| 1 | | 1 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 | | | | | |
| OTC Monograph Drug | M012 | 11/29/2019 | 11/29/2026 | | | | | |

| Marketing Information | | | | | | | | |
|-----------------------|---|-------------------------|-----------------------|--|--|--|--|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | | | | | |
| OTC Monograph Drug | M012 | 11/29/2019 | 11/29/2026 | | | | | |
| | | | | | | | | |

Labeler - CVS Pharmacy (062312574)

Revised: 8/2025 CVS Pharmacy