

**CORICIDIN HBP NIGHTTIME MULTI-SYMPTOM COLD-
acetaminophen, dextromethorphan hydrobromide, doxylamine
succinate liquid
Bayer HealthCare LLC.**

Coricidin HBP Nighttime Multi-Symptom Cold Liquid

Drug Facts

Active ingredients (in each 15 mL)

| | |
|-------------------------------------|-----------------------------|
| Acetaminophen 325 mg | Pain reliever/fever reducer |
| Dextromethorphan hydrobromide 15 mg | Cough suppressant |
| Doxylamine succinate 6.25 mg | Antihistamine |

Purpose

| | |
|-------------------------------------|-----------------------------|
| Acetaminophen 325 mg | Pain reliever/fever reducer |
| Dextromethorphan hydrobromide 15 mg | Cough suppressant |
| Doxylamine succinate 6.25 mg | Antihistamine |

Uses

temporarily relieves common cold/flu symptoms:

- headache
- sore throat
- runny nose
- minor aches and pains
- cough due to minor throat and bronchial irritation
- sneezing
- fever

Warnings

Liver warning

This product contains acetaminophen. Severe liver damage may occur if you take

- more than 120 mL (4 doses) in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Allergy alert

acetaminophen may cause severe skin or severe allergic reactions. Symptoms may

include:

- skin reddening
- blisters
- rash
- hives
- facial swelling
- asthma (wheezing)
- shock

If a skin or general allergic 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

- with any other products 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

- liver disease
- cough that lasts or is chronic such as occurs with smoking, asthma or emphysema
- trouble urinating due to an enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis
- cough that occurs with too much phlegm (mucus)
- glaucoma
- a sodium-restricted diet

Ask a doctor or pharmacist before use if you are

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers

When using this product

- excitability may occur, especially in children
- marked drowsiness may occur
- avoid alcoholic drinks
- be careful when driving a motor vehicle or operating machinery
- alcohol, sedatives, and tranquilizers may increase drowsiness

Stop use and ask a doctor if

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

If pregnant or breast-feeding

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children

Keep out of reach of children. In case of overdose, get medical help

or contact a Poison Control Center right away. (1-800-222-1222) Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

- do not take more than directed
- adults and children 12 years and over: take 30 mL in the dosing cup provided every 6 hours while symptoms last
- do not exceed 120 mL (4 doses) in 24 hours, unless directed by a doctor
- children under 12 years: ask a doctor

Other information

- **each 15 mL contains:** sodium 18 mg
- store at 20°-25°C (68°-77°F)

Inactive ingredients

alcohol 10%, anhydrous citric acid, FD&C blue no.1, FD&C red no.40, flavor, high fructose corn syrup, polyethylene glycol, propylene glycol, purified water, saccharin sodium, sodium citrate

Questions or comments

Questions or comments? 1-800-317-2165 (Mon – Fri 9AM – 5PM EST)

Drug Facts (continued)

Ask a doctor before use if you have

- liver disease
- cough that lasts or is chronic such as occurs with smoking, asthma or emphysema
- trouble urinating due to an enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis
- cough that occurs with too much phlegm (mucus)
- glaucoma ■ a sodium-restricted diet

Ask a doctor or pharmacist before use if you are

- taking the blood thinning drug warfarin ■ taking sedatives or tranquilizers

When using this product

- excitability may occur, especially in children
- marked drowsiness may occur ■ avoid alcoholic drinks
- be careful when driving a motor vehicle or operating machinery
- alcohol, sedatives and tranquilizers may increase drowsiness

Stop use and ask a doctor if

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

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. (1-800-222-1222) Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

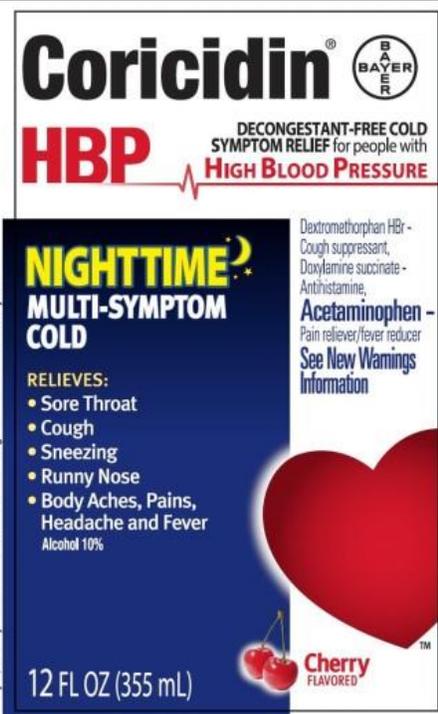
Directions

- do not use more than directed
- adults and children 12 years and over: take 30 mL in the dosing cup provided every 6 hours while symptoms last
- do not exceed 120 mL (4 doses) in 24 hours, unless directed by a doctor
- children under 12: ask a doctor

Other information ■ each 15 mL contains: sodium 18 mg ■ store at 20°-25°C (68°-77°F)

Inactive ingredients alcohol 10%, anhydrous citric acid, FD&C blue no. 1, FD&C red no. 40, flavor, high fructose corn syrup, polyethylene glycol, propylene glycol, purified water, saccharin sodium, sodium citrate

Questions or comments? 1-800-317-2165 (Mon-Fri 9AM - 5PM EST)



Coricidin[®] 

HBP DECONGESTANT-FREE COLD SYMPTOM RELIEF for people with **HIGH BLOOD PRESSURE**

NIGHTTIME[®]
MULTI-SYMPTOM COLD

RELIEVES:

- Sore Throat
- Cough
- Sneezing
- Runny Nose
- Body Aches, Pains, Headache and Fever

Alcohol 10%

12 FL OZ (355 mL)

Cherry FLAVORED

Dextromethorphan HBr - Cough suppressant, Doxylamine succinate - Antihistamine, Acetaminophen - Pain reliever/fever reducer

See New Warnings Information

DO NOT USE IF PRINTED NECKBAND IS BROKEN OR MISSING

PARENTS: Learn about teen medicine abuse www.StopMedicineAbuse.org

DECONGESTANT-FREE COLD SYMPTOM RELIEF for people with HIGH BLOOD PRESSURE

Drug Facts

Active ingredients (in each 15 mL)

| Active ingredients (in each 15 mL) | Purpose |
|-------------------------------------|-----------------------------|
| Acetaminophen 325 mg | Pain reliever/fever reducer |
| Dextromethorphan hydrobromide 15 mg | Cough suppressant |
| Doxylamine succinate 6.25 mg | Antihistamine |

Uses ■ temporarily relieves common cold/flu symptoms:
 ■ headache ■ sore throat ■ runny nose ■ minor aches and pains
 ■ cough due to minor throat and bronchial irritation ■ sneezing ■ fever

Warnings
Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take
 ■ more than 120 mL (4 doses) in 24 hours, which is the maximum daily amount
 ■ with other drugs containing acetaminophen
 ■ 3 or more alcoholic drinks every day while using this product
Allergy alert: acetaminophen may cause severe skin or severe allergic reactions. Symptoms may include:
 ■ skin redness ■ blisters ■ rash ■ hives
 ■ facial swelling ■ asthma (wheezing) ■ shock
 If a skin or general allergic 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
 ■ 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.

CORICIDIN HBP NIGHTTIME MULTI-SYMPTOM COLD

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate liquid

| Product Information | | | |
|--------------------------------|----------------|---------------------------|----------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:11523-4358 |
| Route of Administration | ORAL | | |

| Active Ingredient/Active Moiety | | | |
|--|-------------------------------|------------------|--|
| Ingredient Name | Basis of Strength | Strength | |
| DOXYLAMINE SUCCINATE (UNII: V9B19B5Y12) (DOXYLAMINE - UNII:95QB77JKPL) | DOXYLAMINE SUCCINATE | 6.25 mg in 15 mL | |
| ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D) | ACETAMINOPHEN | 325 mg in 15 mL | |
| DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 15 mg in 15 mL | |

| Inactive Ingredients | | |
|--|----------|--|
| Ingredient Name | Strength | |
| ANHYDROUS CITRIC ACID (UNII: XF417D3PSL) | | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | | |
| FD&C RED NO. 40 (UNII: WZB9127XOA) | | |
| HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S) | | |
| POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) | | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | | |
| WATER (UNII: 059QF0KO0R) | | |
| SACCHARIN SODIUM (UNII: SB8ZUX40TY) | | |

Product Characteristics

| | | | |
|-----------------|--------|---------------------|--|
| Color | | Score | |
| Shape | | Size | |
| Flavor | CHERRY | Imprint Code | |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:11523-4358-1 | 355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 11/20/2015 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M012 | 11/20/2015 | |

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 12/2025

Bayer HealthCare LLC.