NIGHTTIME COUGH- dextromethorphan hydrobromide, doxylamine succinate solution CVS Pharmacy

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

CVS Pharmacy, Inc. Nighttime Cough Drug Facts

Active ingredients (in each 30 mL)

Dextromethorphan HBr 30 mg
Doxylamine succinate 12.5 mg

Purpose

Cough suppressant

Antihistamine

Uses

temporarily relieves cold symptoms:

- · cough due to minor throat and bronchial irritation
- runny nose and sneezing

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

- glaucoma
- cough that occurs with too much phlegm (mucus)
- a breathing problem such as emphysema or chronic bronchitis
- · persistent or chronic cough as occurs with smoking, asthma, or emphysema
- trouble urinating due to an enlarged prostate gland

Ask a doctor or pharmacist before use if you are

When using this product

- excitability may occur, especially in children
- may cause marked drowsiness
- 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

cough persists for more than 1 week, tends to recur, or is accompanied by fever, rash, or persistent headache. A persistent cough may be a sign 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 (1-800-222-1222).

Directions

- take only as directed
- only use the dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over	30 mL every 6 hrs
children 4 to under 12 yrs	ask a doctor
children under 4 yrs	do not use

Other information

- each 30 mL contains: sodium 32 mg
- store at 20-25°C (68-77°F)

Inactive ingredients

alcohol, 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?

1-800-719-9260

Principal Display Panel

CVS Health®

Nighttime Cough

DEXTROMETHORPHAN HBr - Cough suppressant

DOXYLAMINE SUCCINATE - Antihistamine

Relieves: Cough, Sneezing, Runny nose

COUGH

Cherry Flavor

ALCOHOL 10%

12 FL OZ (355 mL)



NDC 59779-668-40

Nighttime Cough

DEXTROMETHORPHAN HBr-Cough suppressant DOXYLAMINE SUCCINATE-Antihistamine

Relieves: Cough, Sneezing, Runny nose

COUGH



Cherry Flavor **ALCOHOL 10%**

12 FL 0Z (355 mL)

: 66840 17 F7

Drug Facts

(continued)

Ask a doctor before use if you

■ a breathing problem such as

persistent or chronic cough as

Ask a doctor or pharmacist before use if you are taking

sedatives or tranquilizers

When using this product

■ avoid alcoholic drinks

excitability may occur, especially

may cause marked drowsiness

■ be careful when driving a motor

vehicle or operating made

cough that occurs with too much

emphysema or chronic bronchitis

occurs with smoking, asthma, or

have ■glaucoma

phleam (mucus)

emphysema ■ trouble urinating due to an enlarged prostate gland

in children

PARENTS:

www.StopMedicineAbuse.org

DO NOT USE IF PRINTED NECKBAND IS BROKEN OR MISSING

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Drug Facts

Active ingredients (in each 30 mL)

Dextromethorphan HBr 30 mg......Cough suppressant

Purpose

Doxylamine succinate 12.5 mg......Antihistamine

Uses temporarily relieves cold symptoms:

- cough due to minor throat and bronchial irritation
- runny nose and sneezing

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.

PEEL BACK AT CORNER FOR MORE INFORMATION

: 66840 17 81

Drug Facts (continued)

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).

Directions

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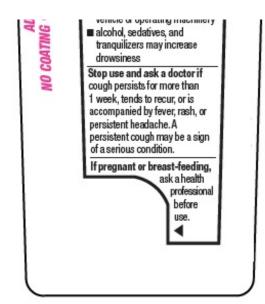
Inactive ingredients

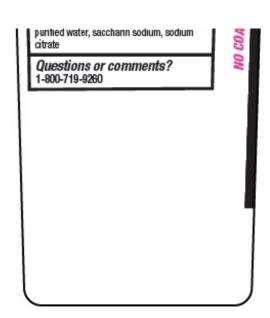
alcohol, anhydrous citric acid, FD&C blue no. 1, FD&C red no. 40, flavor, high fructose corn syrup, polyethylene glycol, propylene glycol,

NO VARNISH . NO TYPE

■ store at 20-25°C (68-77°F)

• NO TYPE





NIGHTTIME COUGH

dextromethorphan hydrobromide, doxylamine succinate solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-668
Route of Administration ORAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 30 mL		
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 30 mL		

Inactive Ingredients				
Ingredient Name	Strength			
ALCOHOL (UNII: 3K9958V90M)				
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)				
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)				

Product Characteristics			
Color	RED (Dark Red)	Score	

Shape	Size	
Flavor	Imprint Code	
Contains		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		295 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/23/2002	11/09/2013
2	NDC:59779-668- 40	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/24/2008	

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
OTC monograph final	part341	09/23/2002	

Labeler - CVS Pharmacy (062312574)

Revised: 12/2022 CVS Pharmacy