

TUSSIN DM DAYTIME NIGHTTIME- dextromethorphan hbr, doxylamine succinate, guaifenesin
CVS WOONSOCKET PRESCRIPTION CENTER, INCORPORATED

CVS 44-030043

Day

Active ingredients (in each 20 mL)

Dextromethorphan HBr 20 mg
Guaifenesin 400 mg

Purpose

Cough suppressant
Expectorant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

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

- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema

Stop use and ask a doctor if

cough persists more than 7 days, tends to recur, or is accompanied by a 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

- **do not take more than directed**
- do not take more than 6 doses in any 24-hour period
- mL = milliliter
- only use the dose cup provided
- adults and children 12 years and over: 20 mL in dosing cup provided every 4 hours
- children under 12 years: do not use

Other information

- **each 20 mL contains:** sodium 16 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

anhydrous citric acid, FD&C blue #1, FD&C red #40, flavors, glycerin, high fructose corn syrup, microcrystalline cellulose, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium chloride, sodium citrate dihydrate, sorbitol, sucralose, xanthan gum

Questions or comments?

1-800-426-9391

Night

Active ingredients (in each 20 mL)

Dextromethorphan HBr 30 mg
Doxylamine Succinate 12.5 mg

Purpose

Cough suppressant
Antihistamine

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
 - itching of the nose or throat
 - itchy, watery eyes
 - runny nose
 - sneezing
- controls the impulse to cough to help you 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

- a cough that occurs with too much phlegm (mucus)
- glaucoma
- difficulty in urination due to enlargement of the prostate gland
- a breathing problem or persistent or chronic cough as occurs with smoking, asthma, chronic bronchitis, or emphysema

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

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

Stop use and ask a doctor if

cough persists more than 7 days, tends to recur, or is accompanied by a 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

- **do not take more than directed**
- do not take more than 4 doses in any 24-hour period
- mL = milliliter
- only use the dose cup provided
- adults and children 12 years and over: 20 mL in dosing cup provided every 6 hours
- children under 12 years: do not use

Other information

- **each 20 mL contains:** sodium 14 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

anhydrous citric acid, FD&C blue #1, FD&C red #40, flavors, glycerin, high fructose corn syrup, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium chloride, sodium citrate dihydrate, sucralose, sugar, xanthan gum

Principal Display Panel

DAY & NIGHT COMBO PACK

<p>♥CVS Health® Compare to the active ingredients in Robitussin® Maximum Strength Cough + Chest Congestion DM* MAXIMUM STRENGTH MAXIMUM STRENGTH FOR MUCUS RELIEF Daytime Non-Drowsy Tussin DM DEXTROMETHORPHAN HBr Cough suppressant GUAIFENESIN Expectorant Cough & Chest Congestion Relieves:</p> <ul style="list-style-type: none">• Cough• Chest congestion• Mucus <p>Menthol-Berry Flavor Dosage cup provided For Ages 12 & Over Actual Bottle Size on Side Panel 4 FL OZ (118 mL) + 4 FL OZ (118 mL) TOTAL 8 FL OZ (236 mL)</p>	<p>♥CVS Health® Compare to the active ingredients in Robitussin® Maximum Strength Nighttime Cough DM* MAXIMUM STRENGTH MAXIMUM STRENGTH Nighttime Tussin DM DEXTROMETHORPHAN HBr Cough suppressant DOXYLAMINE SUCCINATE Antihistamine Cough & Antihistamine Relieves:</p> <ul style="list-style-type: none">• Cough• Runny nose• Sneezing <p>Menthol-Berry Flavor Dosage cup provided For Ages 12 & Over Actual Bottle Size on Side Panel</p>
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DO NOT TAKE DAYTIME AND NIGHTTIME PRODUCTS AT THE SAME TIME

TAMPER EVIDENT: DO NOT USE IF PRINTED NECK WRAP IS BROKEN OR MISSING

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

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guaranteed.**

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PARENTS:

Learn about teen medicine abuse

www.StopMedicineAbuse.org

Package Contains Two Bottles

Actual Size

Tussin DM
Daytime Non-Drowsy
FOR MUCUS RELIEF
MAXIMUM STRENGTH

Tussin DM
Nighttime
MAXIMUM STRENGTH

CVS Health

DAY & NIGHT COMBO PACK

CVS Health **CVS Health**

MAXIMUM STRENGTH **MAXIMUM STRENGTH**

Tussin DM
Daytime Non-Drowsy
Tussin DM
Nighttime

DEXTROMETHORPHAN HBR
Cough suppressant
QUAIFENESIN
Expectorant
Cough & Chest Congestion

Relieves:
• Cough
• Chest congestion
• Mucus

Menthol-Berry Flavor
Dosage cup provided

Menthol-Berry Flavor
Dosage cup provided

4 FL OZ (118 mL) + 4 FL OZ (118 mL)
TOTAL 8 FL OZ (236 mL)

Daytime, Tussin DM (continued)

Drug Facts (continued)

Other information

Warnings

Directions

Other information

Inactive ingredients

#181525

FPO 100%
UPC# 050428393444

X XXXXXX XXXXXX X

No Print / No Varnish
Lot no. & Exp. date

B-0231-030/043-36H
ORG012303004336

CVS 44-030043

TUSSIN DM DAYTIME NIGHTTIME

dextromethorphan hbr, doxylamine succinate, guaifenesin kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51316-434
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51316-434-19	1 in 1 PACKAGE, COMBINATION; Type 0: Not a Combination Product	04/18/2023	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE, PLASTIC	118 mL
Part 2	1 BOTTLE, PLASTIC	118 mL

Part 1 of 2

TUSSIN DM DAYTIME

dextromethorphan hbr, guaifenesin solution

Product Information

Item Code (Source) NDC:51316-304

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)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	red (MAROON)	Score	
Shape		Size	
Flavor	MENTHOL, BERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		118 mL in 1 BOTTLE, PLASTIC; 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	04/18/2023	

Part 2 of 2

TUSSIN DM NIGHTTIME

dextromethorphan hbr, doxylamine succinate solution

Product Information

Item Code (Source)	NDC:51316-430
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 20 mL
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 20 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)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SUCROSE (UNII: C151H8M554)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	red (MAROON)	Score	
Shape		Size	
Flavor	MENTHOL, BERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		118 mL in 1 BOTTLE, PLASTIC; 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	04/18/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	04/18/2023	

Labeler - CVS WOONSOCKET PRESCRIPTION CENTER, INCORPORATED (062312574)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(51316-434) , pack(51316-434)

Revised: 8/2025

CVS WOONSOCKET PRESCRIPTION CENTER, INCORPORATED