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

- do not exceed recommended dosage
- 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

VCVS

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:
• Cough

Chest congestion

Mucus

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)

♥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:

Cough

Runny nose

Sneezing

Menthol-Berry Flavor

Dosage cup provided For Ages 12 & Over

Actual Bottle Size on Side Panel

50844 ORG012303004336

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

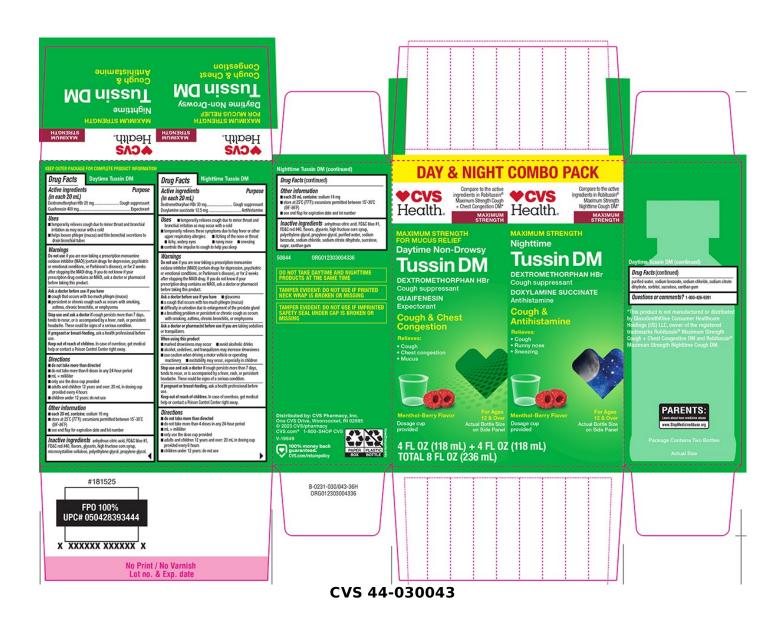
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PARENTS:

Learn about teen medicine abuse www.StopMedicineAbuse.org

Package Contains Two Bottles Actual Size



TUSSIN DM DAYTIME NIGHTTIME

dextromethorphan hbr, doxylamine succinate, quaifenesin kit

Product Information

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

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

4	(
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

#	ltem 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		

Strength
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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: 8/2023 CVS WOONSOCKET PRESCRIPTION CENTER, INCORPORATED