WALGREENS DAY AND NIGHT PACK- dextromethorphan hbr, guaifenesin, chlorpheniramine maleate
WALGREEN CO

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.

Walgreen Children's Daytime & Night-Time Cough and Chest Congestion DM

Children's Daytime Wal-Tussin® DM

Active ingredients Day Time (in each 20 mL)

Dextromethorphan HBr USP 20 mg

Guaifenesin. USP 200 mg

Children's Nighttime Wal-Tussin® DM

Active ingredients for Nighttime (in each 10 mL)

Chlorpheniramine maleate, USP 2 mg

Dextromethorphan HBr USP 15 mg

Purposes for Daytime Wal-Tussin® DM

Cough suppressant

Expectorant

Purpose for Nighttime Wal-Tussin® DM

Antihistamine

Cough suppressant

Uses

DAYTIME

- 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

NIGHTTIME

- 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:
    - runny nose
    - sneezing
    - itching of the nose or throat

- itchy, watery eyes

Warnings

Do not us

DAYTIME

- 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.

NIGHTTIME

- to sedate a child or to make a child sleepy
- 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

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

NIGHTTIME

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

Ask a doctor or pharmacist before use if you are

NIGHTTIME

taking sedatives or tranquilizers.

When using this product

NIGHTTIME

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

Stop use and ask a doctor if

DAYTIME

- cough last more than 7 days, comes back or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition.

NIGHTTIME

- cough last more than 7 days, comes back or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition.

If pregnant or breast-feeding,

DAYTIME

ask a health professional before use.

NIGHTTIME

ask a health professional before use

Keep out of reach of children.

DAYTIME

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

NIGHTTIME

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away

Directions

Daytime Wal-Tussin® DM

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided.
- keep dosing cup with product
- mL= milliliter

age	dose
Children under 4 years	Do not use
Children 4 to under 6 years	5 mL every 4 hours
Children 6 to under 12 years	10 mL every 4 hours
Adults and children 12 years and older	20 mL every 4 hours

Nighttime Wal-Tussin® DM

- do not take more than 4 doses in any 24-hour period
- measure only with dosing cup provided.
- keep dosing cup with product
- mL= milliliter

age	dose

Children under 6 years	Do not use
Children 6 to under 12 years	10 mL every 6 hours
Adults and children 12 years and older	20 mL every 6 hours

Other information

DAYTIME Wal-Tussin DM

- **each 20 mL contains:** sodium 11 mg
- store at room temperature. Do not refrigerate
- contain low sodium
- **do not use if printed seal under cap is torn or missing**

Nighttime Wal-Tussin DM

- **each 10 mL contains:** sodium 6 mg
- store at room temperature. Do not refrigerate
- contain low sodium
- **do not use if printed seal under cap is torn or missing**

Inactive ingredients

***Inactive ingredients for Day Time* Wal-Tussin DM**

anhydrous citric acid, carboxymethylcellulose sodium, edetate disodium, FD &C Blue No. 1, FD&C Red No. 40, natural and artificial flavors, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum

***Inactive ingredients for Night Time* Wal-Tussin DM**

anhydrous citric acid, FD&C Red No. 40, natural and artificial flavors, glycerin, lactic acid, potassium sorbate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose

Questions or comments?

1-866-467-2748

Principal Display Panel

Compare to Children's Robitussin® Cough & Chest Congestion DM* & Nighttime Cough DM Long-Active active ingredients††

Children's

Cough & Chest Congestions DM

Wal-Tussin® DM

DEXTROMETHORPHAN HBr, USP 20 mg/ 20 mL

COUGH SUPPRESSANT

GUAIFENESIN, USP 200 mg

EXPECTORANT

DAYTIME

NON-DROWSY

Relieves cough, chest congestion & mucus

4 YEARS & OVER

GRAPE FLAVOR

Naturally and Artificially Flavored

Dosage cup included

2 - 4 FL OZ (118 mL) BOTTLES / TOTAL – 8 FL OZ (236 mL)

††This product is not manufactured or distributed by Pfizer, the distributor of Children's Robitussin® Cough & Chest Congestion DM.

Children's

Nighttime Cough DM

Wal-Tussin® DM

CHLORPHENIRAMINE MALEATE, USP 2 mg / 10 mL

ANTIHISTAMINE

DEXTROMETHORPHAN HBr, USP 15 mg / 10 mL

COUGH SUPPRESSANT

NIGHTTIME

- Relieves cough & runny nose

Alcohol free

6 YEARS & OLDER

Fruit Punch Flavor

Naturally and Artificially Flavored

Dosage cup included

2 - 4 FL OZ (118 mL) BOTTLES TOTAL – 8 FL OZ (236 mL)

††This product is not manufactured or distributed by Pfizer, the distributor of Children's Robitussin® Nighttime Cough DM Long-Acting.

Walgreens

PHARMACIST RECOMMENDED

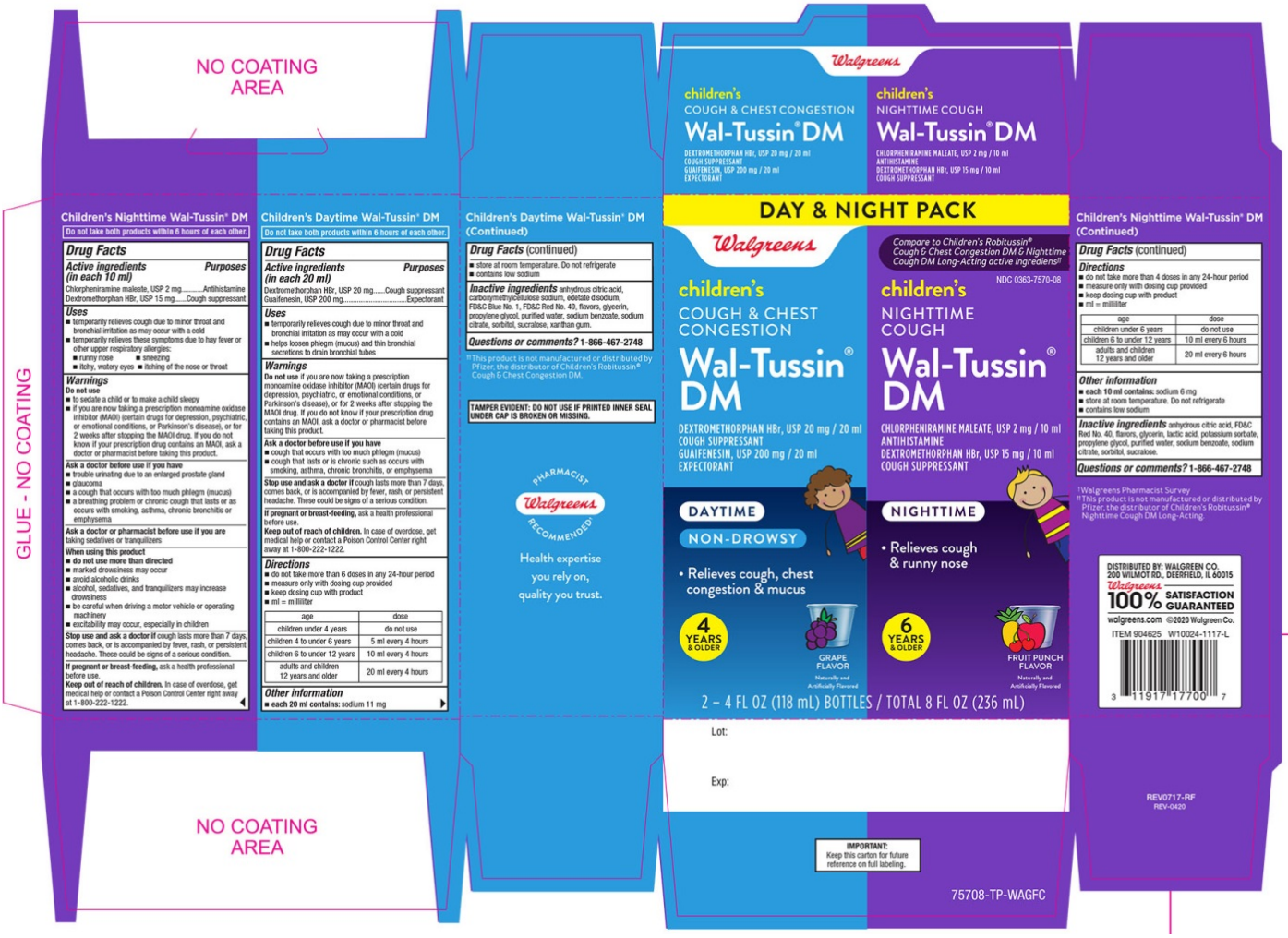
TAMPER EVIDENT: DO NOT USE IF PRINTED INNER SEAL UNDER CAP IS BROKEN OR MISSING

DISTRIBUTED BY: WALGREENS CO.

200 WILMOT RD. DEERFIELD, IL 60015

Walgreens 10% SATISFACTION GUARANTEED

walgreens.com ©2020 Walgreen Co.



WALGREENS DAY AND NIGHT PACK

dextromethorphan hbr, guaifenesin, chlorpheniramine maleate kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-7570

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-7570-08	1 in 1 KIT; Type 9: Other Type of Part 3 Combination Product (e.g., Drug/Device/Biological Product)	04/13/2020	

Quantity of Parts

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

Part 1 of 2

DAYTIME COUGH AND CHEST CONGESTION

dextromethorphan hbr, guaifenesin liquid

Product Information

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	200 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	PURPLE	**Score**	
Shape		**Size**	
Flavor	GRAPE	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		118 mL in 1 BOTTLE; Type 9: Other Type of Part 3 Combination Product (e.g., Drug/Device/Biological Product)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	04/13/2020	

Part 2 of 2

NIGHT TIME COUGH DM

chlorpheniramine maleate, diphenhydramine hbr liquid

Product Information

Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg in 10 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
LACTIC ACID, UNSPECIFIED FORM (UNII: 33X04XA5AT)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color	RED	**Score**	
Shape		**Size**	
Flavor	FRUIT PUNCH	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1		118 mL in 1 BOTTLE; Type 9: Other Type of Part 3 Combination Product (e.g., Drug/Device/Biological Product)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	04/13/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	04/13/2020	

Labeler - WALGREEN CO (008965063)

Revised: 5/2020 WALGREEN CO