

**CHILDRENS COUGH AND CONGESTION, CHILDRENS COLD AND COUGH DAY,
NIGHT- dextromethorphan hbr, doxylamine succinate, guaifenesin
WALMART INC.**

Equate 44-019020

Day Children's Cough & Congestion

Active ingredients (in each 15 mL)

Dextromethorphan HBr 10 mg
Guaifenesin 100 mg

Purpose

Cough suppressant
Expectorant

Uses

- temporarily relieves these common cold symptoms:
 - cough due to minor throat and bronchial irritation
 - cough to help you sleep
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

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

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

Stop use and ask a doctor if

cough persists more than 1 week, 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 exceed 6 doses per 24 hours
- mL = milliliter
- only use the dose cup provided

adults and children 12 years and over	30 mL every 4 hours
children 6 to under 12 years	15 mL every 4 hours
children under 6 years	do not use

Other information

- **each 15 mL contains:** sodium 8 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, D&C yellow #10, FD&C green #3, FD&C red #40, FD&C yellow #6, flavor, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sodium saccharin, sorbitol solution, sucralose, xanthan gum

Questions or comments?

1-888-287-1915

Night Children's Cold & Cough

Active ingredients (in each 15 mL)

Dextromethorphan HBr 10 mg
Doxylamine succinate 6.25 mg

Purpose

Cough suppressant
Antihistamine

Uses

temporarily relieves these common cold symptoms:

- cough due to minor throat and bronchial irritation
- cough to help you sleep
- 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.
- to make a child sleepy

Ask a doctor before use if you have

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

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

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

Stop use and ask a doctor if

cough persists more than 1 week, 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 exceed 6 doses per 24 hours
- mL = milliliter
- only use the dose cup provided

adults and children 12 years and over	30 mL every 4 hours
children 6 to under 12 years	15 mL every 4 hours

children under 6 years	do not use
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Other information

- **each 15 mL contains:** sodium 8 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, D&C yellow #10, FD&C green #3, FD&C red #40, FD&C yellow #6, flavor, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sodium saccharin, sorbitol solution, sucralose, xanthan gum

Principal display panel

DAY & NIGHT VALUE PACK

HONEY
FLAVORED

NDC 79903-414-16

equate™ children's Cough & Congestion Day Ages 6+ Day Cough & Congestion Dextromethorphan HBr (Cough Suppressant) Guaifenesin (Expectorant) Relieves: <ul style="list-style-type: none">• Chest congestion• Cough 8 FL OZ (237 mL) Day 16 FL OZ (474 mL) TOTAL	equate™ children's Cold & Cough Night Ages 6+ Night Cold & Cough Dextromethorphan HBr (Cough Suppressant) Doxylamine succinate (Antihistamine) Relieves: <ul style="list-style-type: none">• Sneezing• Runny nose• Cough 8 FL OZ (237 mL) Night
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**TAMPER EVIDENT: DO NOT USE IF
PRINTED NECK WRAP IS BROKEN OR
MISSING**

**Do Not Take Daytime and Nighttime
Products at the Same Time.**

PARENTS:

Learn about teen medicine abuse
www.StopMedicineAbuse.org

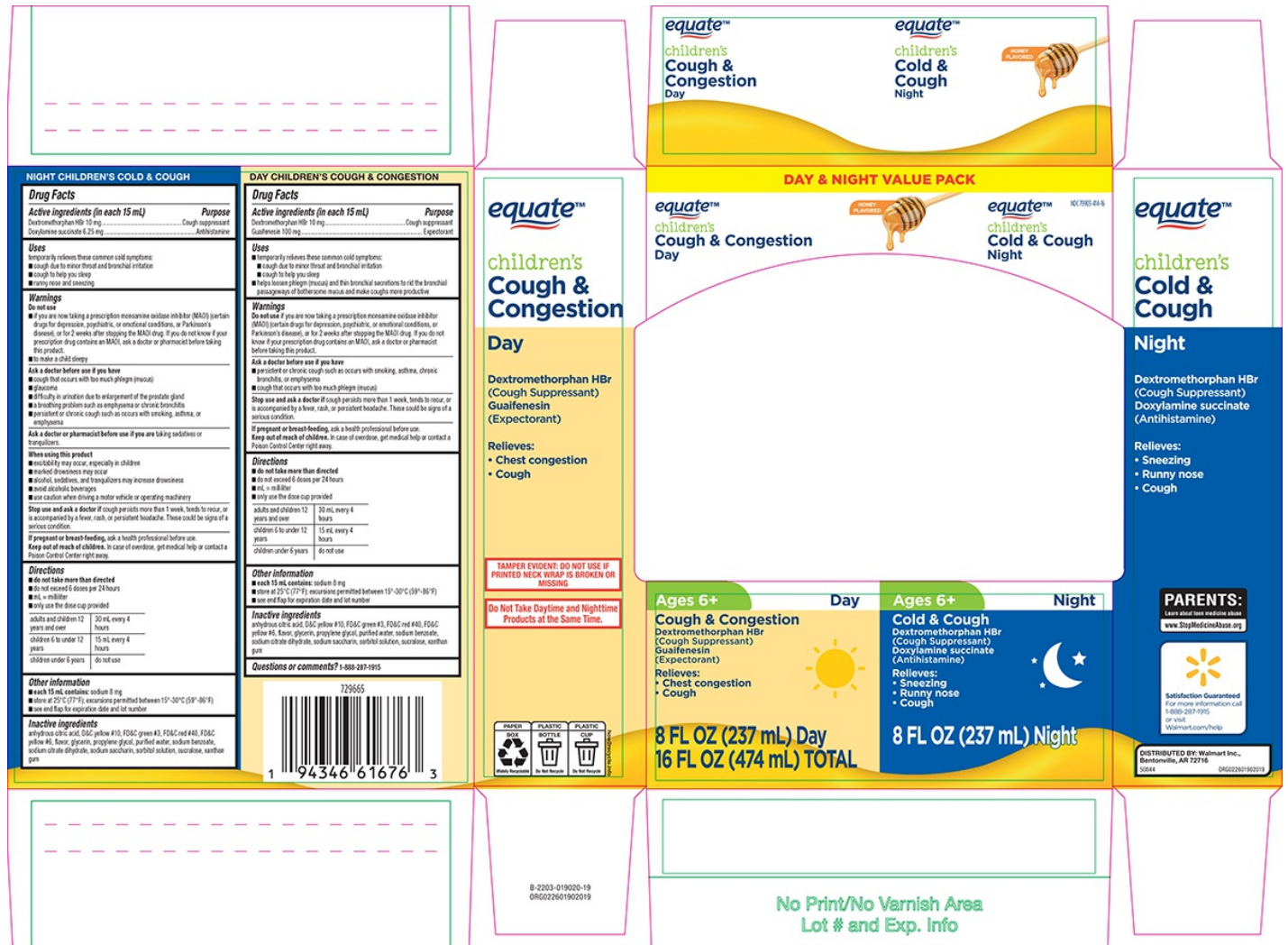
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For more information call
 1-888-287-1915
 or visit
Walmart.com/help

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ORG022601902019



Equate 44-019020

CHILDRENS COUGH AND CONGESTION, CHILDRENS COLD AND COUGH DAY, NIGHT

dextromethorphan hbr, doxylamine succinate, guaifenesin kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-414
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-414-16	1 in 1 PACKAGE	05/19/2026	
1		1 in 1 KIT; Type 0: Not a Combination Product		

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE, PLASTIC	237 mL
Part 2	1 BOTTLE, PLASTIC	237 mL

Part 1 of 2

CHILDRENS COUGH AND CONGESTION DAY

dextromethorphan hbr, guaifenesin solution

Product Information

Item Code (Source)	NDC:79903-412
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 15 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	

SODIUM SACCHARIN (UNII: SB8ZUX40TY)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	brown (light brown)	Score	
Shape		Size	
Flavor	HONEY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-412-08	237 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	05/19/2026	

Part 2 of 2

CHILDRENS COLD AND COUGH NIGHT

dextromethorphan hbr, doxylamine succinate solution

Product Information

Item Code (Source)	NDC:79903-413
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 15 mL
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)
FD&C RED NO. 40 (UNII: WZB9127XOA)
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)
GLYCERIN (UNII: PDC6A3C0OX)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
WATER (UNII: 059QF0KO0R)
SODIUM BENZOATE (UNII: OJ245FE5EU)
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)
SODIUM SACCHARIN (UNII: SB8ZUX40TY)
SORBITOL SOLUTION (UNII: 8KW3E207O2)
SUCRALOSE (UNII: 96K6UQ3ZD4)
XANTHAN GUM (UNII: TTV12P4NEE)

Product Characteristics

Color	brown (light brown)	Score	
Shape		Size	
Flavor	HONEY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-413-08	237 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	05/19/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	05/19/2026	

Labeler - WALMART INC. (051957769)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(79903-414) , pack(79903-414)

