WALGREENS NIGHTTIME KIDS HONEY COLD COUGH CONGESTION- doxylamine succinate, phenylephrine hydrochloride and dextromethorphan hydrobromide liquid WALGREENS CO

Walgreens Nighttime Kids Cold & Cough + Congestion

Drug Facts

Active ingredients (in each 15 mL)

Dextromethorphan HBr 10 mg
Doxylamine succinate 6.25 mg
Phenylephrine HCl 5 mg

Purpose

Cough suppressant Antihistamine Nasal decongestant

Uses

temporarily relieves common cold symptoms:

- nasal congestion
- sinus congestion & pressure
- cough due to minor throat & bronchial irritation
- cough to help you sleep
- runny nose & sneezing
- reduces swelling of nasal passages
- temporarily restores freer breathing through the nose
- promotes nasal and/or sinus drainage

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

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- glaucoma
- trouble urinating due to enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts or 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

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

Stop use and ask a doctor if

- you get nervous, dizzy or sleepless
- symptoms do not improve within 7 days or occur with a fever
- cough persists for more than 7 days, comes back or occurs with 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.

Overdose warning

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

Directions

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

adults & children 12 yrs & over	30 mL every 4 hrs
children 6 to under 12 yrs	15 mL every 4 hrs
children 4 to under 6 yrs	do not use unless directed by a doctor

Other information

- each 15 mL contains: sodium 12 mg
- store at room temperature

Inactive ingredients

Anhydrous citric acid, carboxymethylcellulose sodium, disodium edetate, D&C Yellow No. 10, FD&C Green No. 3, FD&C Red No. 40, FD&C Yellow No. 6, flavor, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose, , xanthan gum

Questions?

1-866-467-2748

Distributed by:

PRINCIPAL DISPLAY PANEL - 236 ml Bottle Label

Compare to the active ingredients in Kids **Vicks**[®] NyQuil[™] Honey Cold & Cough + Congestion*

NDC 0363-6248-08

Nighttime

Kid's

Cold & Cough + Congestion

Doxylamine Succinate, Phenylephrine HCl, Dextromethorphan HBr

- Sneezing, Runny Nose
- Nasal Congestion
- Cough

No Added Alcohol & Acetaminophen Free

Honey Flavor

Naturally and Artificially Flavored

8 FL OZ (236 mL)

*This Product is not manufactured or distributed by Procter & Gamble, the distributor of Kids **Vicks**[®] NyQuil™ Honey Cold & Cough + Congestion.





BACK 6.75"Width

Drug Facts (continued) Pipregnant or breast-feeding, ask a health professional before use Reep out of reach of children. In case of everces, get medical help or contact a Poison Control Center right away at 1-900-222-1222. Directions ■ take only as directed ■ only use the dose out provided ■ do not exceed 4 doses par 24 hrs adults & children 12 yrs & over | | \$0 mL every 4 hrs children 6 to under 12 year 15 mL every 4 hrs. 3.25" Height children 4 to under 6 vis. do not use unless directed by a doctor. children under 4 vrs. do not use Other information ■ each 15 mL contains: sodium 12 mg ■ store at room temperature Inactive ingredients artistics of to add, cartesymetryl solutes and in 1986 Yelva No. 10, dead on obotic FDSC Gree No. 90, FDSC No. 80, FDSC No. 80, FDSC No. 80, FDSC No. 90, PSC No. 80, FDSC NO. 80, Questions or comments? 1-996-467-2748

WALGREENS NIGHTTIME KIDS HONEY COLD COUGH CONGESTION

doxylamine succinate, phenylephrine hydrochloride and dextromethorphan hydrobromide liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-6248
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
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 15 mL

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)		
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311)		
EDETATE DISODIUM (UNII: 7FLD91C86K)		
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)		
PROPYL GALLATE (UNII: 8D4SNN7V92)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SORBITOL (UNII: 506T60A25R)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		
XANTHAN GUM (UNII: TTV12P4NEE)		

Product Characteristics			
Color	brown	Score	
Shape		Size	
Flavor	HONEY	Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363- 6248-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/01/2023	

Labeler - WALGREENS CO (008965063)

Revised: 11/2024 WALGREENS CO