

HONEYWORKS PLUS ADULT NIGHTTIME COUGH SYRUP- dextromethorphan hbr, doxylamine succinate solution
RARITAN PHARMACEUTICALS INC

HoneyWorks adult Cough Syrup Drug facts

<i>Active ingredients (in each 20 ml)</i>	<i>Purposes</i>
Dextromethorphan HBr, USP 20 mg	Cough suppressant
Doxylamine Succinate, USP 12.5 mg	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:
 - runny nose
 - sneezing
 - itchy, watery eyes
 - itching of the nose or throat
- controls the impulse to cough to help you sleep

Warnings

Do not use

- 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

- trouble urinating due to an enlarged 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 taking sedatives or tranquilizers

When using this product

- **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 cough lasts 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, 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

- measure only with dosing cup provided
- keep dosing cup with product
- ml = milliliter
- do not take more than 4 doses in any 24-hour period
- this adult products is not intended for use in children under 12 years of age

age	dose
adults and children 12 years and older	20 ml every 6 hours
children under 12 years	Do not use

Other information

- each 20 ml contains: **sodium 3 mg**
- store at room temperature

Inactive ingredients

citric acid, natural flavor, organic honey, purified water, sodium benzoate,

Questions or comments?

1-866-467-2748

Made in the U.S.A. with domestic and foreign ingredients

Distributed by Lifelab Health LLC

5300 W Hillsboro Blvd, Suite 202,

Coconut Creek, FL 33073

PRINCIPAL DISPLAY PANEL - 118 ml Bottle Carton

NDC 68163-660-08

HONEYWORKS™ Plus

Adult

COUGH SYRUP

Dextromethorphan HBr
(COUGH SUPPRESSANT)

Doxylamine Succinate (ANTIHISTAMINE)

NIGHTTIME FORMULA

THIS GREAT-TASTING PRODUCT OFFERS FAST & EFFECTIVE COUGH RELIEF NAD IS MADE WITH ORGANIC DARK HONEY

- NO PARABENS
- NO DYES
- NO HIGH FRUCTOSE CORN SYRUP
- NO ARTIFICIAL FLAVORS
- NON-GMO
- SOY-FREE
- GLUTEN FREE
- DAIRY-FREE
- PEANUT-FREE
- TREE NUT-FREE
- ZERO TRANS FAT

FOR AGES 12 YEARS & OLDER

ALSO TRY HONEYWORKS™ KIDS SOOTHING THROAT SPRAY

8 FL OZ (236 mL)

IMPORTANT: Keep this carton for future reference on full labeling.



HONEYWORKS PLUS ADULT NIGHTTIME COUGH SYRUP

dextromethorphan hbr, doxylamine succinate solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68163-660
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
DOXYLAMINE SUCCINATE (UNII: V9B19B5Y12) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
HONEY (UNII: Y9H1V576FH)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68163-660-08	1 in 1 CARTON	02/01/2023	
1		236 mL in 1 BOTTLE; 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	02/01/2023	

Labeler - RARITAN PHARMACEUTICALS INC (127602287)

Revised: 10/2025

RARITAN PHARMACEUTICALS INC