

**NIGHT TIME COLD/FLU RELIEF CHERRY - acetaminophen, dextromethorphan hbr, doxylamine succinate liquid**  
**Kareway Product, Inc.**

*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.*

-----

**Drug Facts**

**Active ingredients (in each 15 ml tablespoon)**

Acetaminophen 500 mg

Dextromethorphan HBr 15 mg

Doxylamine succinate 6.25 mg

**Purpose**

Pain reliever/fever reducer

Cough suppressant

Antihistamine

**Uses**

temporarily relieves common cold/flu symptoms:

- runny nose and sneezing
- cough due to minor throat and bronchial irritation
- sore throat
- headache
- minor aches and pains
- fever

**Warnings**

**Liver warning:** This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4 doses in 24 hours which is the maximum daily amount for this product
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks daily while using this product

**Sore throat warning:** If sore throat is severe, persists for more than 2 days, occurs with or is followed by fever, headache, rash, nausea, or vomiting, see a doctor promptly.

**Do not use**

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- 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 sleep

**Ask a doctor before use if you have**

- liver disease
- glaucoma
- 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
- trouble urinating due to enlarged prostate gland

**Ask a doctor or pharmacist before use if you are**

- taking sedatives or tranquilizers.
- taking the blood thinning drug warfarin

**When using this product**

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

**Stop use and ask a doctor if**

- pain or cough gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back or occurs with rash or headache that lasts.

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:**

Taking more than directed can cause serious health problems. In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

**Directions**

- take only as directed - see Overdose warning
- do not exceed 4 doses per 24 hours

|                                     |                              |
|-------------------------------------|------------------------------|
| adults and children 12 yrs and over | 30 ml (2 Tbsp) every 6 hours |
| children 4 to under 12 yrs          | ask a doctor                 |

children under 4 yrs

do not use

### Other information

- each 30 mL dose cup contains: sodium 43 mg
- store at room temperature

### Inactive ingredients

alcohol, citric acid, FD and C blue no.1, FD and C red no.40, flavor, high fructose corn syrup, polyethylene glycol, propylene glycol, purified water, saccharin sodium, sodium citrate, sucrose, sucralose

### Package label

Night Time Cold/Flu Relief Original



## NIGHT TIME COLD/FLU RELIEF CHERRY

acetaminophen, dextromethorphan hbr, doxylamine succinate liquid

### Product Information

|                                |                |                           |                |
|--------------------------------|----------------|---------------------------|----------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:67510-0507 |
| <b>Route of Administration</b> | ORAL           |                           |                |

### Active Ingredient/Active Moiety

| Ingredient Name  | Basis of Strength                | Strength            |
|--|----------------------------------|---------------------|
| <b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)                    | ACETAMINOPHEN                    | 500 mg<br>in 15 mL  |
| <b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN<br>HYDROBROMIDE | 15 mg<br>in 15 mL   |
| <b>DOXYLAMINE SUCCINATE</b> (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)                | DOXYLAMINE SUCCINATE             | 6.25 mg<br>in 15 mL |

## Inactive Ingredients

| Ingredient Name                             | Strength |
|---|----------|
| ALCOHOL (UNII: 3K9958V90M)                  |          |
| CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)  |          |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD)          |          |
| FD&C RED NO. 40 (UNII: WZB9127XOA)          |          |
| HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S) |          |
| POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)     |          |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3)         |          |
| WATER (UNII: 059QF0K00R)                    |          |
| SACCHARIN SODIUM (UNII: SB8ZUX40TY)         |          |
| SODIUM CITRATE (UNII: 1Q73Q2JULR)           |          |
| SUCROSE (UNII: C151H8M554)                  |          |
| SUCRALOSE (UNII: 96K6UQ3ZD4)                |          |

## Packaging

| # | Item Code        | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---------------------|----------------------|--------------------|
| 1 | NDC:67510-0507-4 | 120 mL in 1 BOTTLE  |                      |                    |
| 2 | NDC:67510-0507-6 | 177 mL in 1 BOTTLE  |                      |                    |
| 3 | NDC:67510-0507-0 | 295 mL in 1 BOTTLE  |                      |                    |
| 4 | NDC:67510-0507-2 | 354 mL in 1 BOTTLE  |                      |                    |

## Marketing Information

| Marketing Category  | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---------------------|--|----------------------|--------------------|
| OTC monograph final | part341                                  | 08/26/2011           |                    |

**Labeler** - Kareway Product, Inc. (121840057)

Revised: 8/2011

Kareway Product, Inc.