#### DAYTIME COLD AND FLU- acetaminophen dextromethorphan hbr phenylephrine hcl liquid QUALITY CHOICE (Chain Drug Marketing Association)

\_\_\_\_\_

#### **Drug Facts**

#### Active ingredients (in each 15 mL)

#### Acetaminophen 325 mg

Dextromethoprhan HBr 10 mg

Phenlyephrine HCl 5 mg

## Purposes

#### Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

#### Uses

- temporarily relieves these common cold/flu symptoms:
  - minor aches and pains
  - headache
  - sore throat
  - nasal congestion
  - fever
  - cough due to minor throat and bronchial irritation

## Warnings

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

- adult takes more than 4 doses (30 mL each) of acetaminophen in 24 hours, which is the maximum daily amount
- child takes more than 4 doses (15 mL each) in 24 hours
- taken with other drugs containing acetaminophen
- adult has 3 or more alcoholic drinks every day while using this product

Allery alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away

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

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

# Ask a doctor before use if you have

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- a sodium-restricted diet
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking, asthma or emphysema
- cough that occurs with too much phlegm (mucus)

# Ask a doctor or pharmacist before use if

you are taking the blood thinning drug warfarin.

# When using this product,

# do not exceed recommended dosage.

## Stop use and ask a doctor if

- nervousness, dizziness or sleeplessness occurs
- pain, nasal congestion or cough gets worse or lasts more than 5 days (children) or 7 days (adults)
- 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 the recommended dose (overdose) may cause

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

- do not take more than directed (see Overdose warning)
- do not exceed 4 doses in any 24 hour-period
- measure only with dosing cup provided. Do not use any other dosing device
- keep dosing cup with product
- mL=milliliter

| adults and children 12 years and over | 30 mL every 4 hours |
|---------------------------------------|---------------------|
| children 6 to under 12 years          | 15 mL every 4 hours |
| children 4 to under 6 years           | ask a doctor        |
| children under 4 years                | do not use          |

• When using Daytime and Nighttime products, carefully read each label to ensure correct dosing

#### Other information

- each 15 mL contains: sodium 12 mg
- store between 20-25°C (68-77°F). Do not refrigerate

## Inactive ingredients

citric acid, FD&C yellow #6, flavor, glycerin, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol, sucralose, xanthan gum

#### **Questions or comments?**

Call

## **Principal Display Panel**

\*Compare to the active ingredients in Vicks® DayQuil® Cold & Flu

Non-Drowsy

Multi-Symptom

## Daytime

Cold & Flu Relief

## Acetaminophen 325 mg

Pain reliever/Fever reducer

Dextromethorphan HBr 10 mg

Cough suppressant

Phenylephrine HCl 5 mg

Nasal decongestant

Relieves:

- Aches, fever & sore throat
- Cough
- Nasal congestion

For ages 6 years and over

Alcohol-free

Antihistamine-free

FL OZ (mL)

## TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL AROUND DOSAGE CUP OR UNDER CAP IS BROKEN OR MISSING.

\*This product is not manufactured or distributed by The Procter & Gamble Company. Vicks® and DayQuil® are registered trademarks of The Procter & Gamble Company.

Manufactured by:

Package Label

NDC XXXXX-XXX-XX

\*Compare to the active ingredients in Vicks® DayQuit® Cold & Flu

Non-Drowsy Multi-Symptom

# Daytime Cold & Flu Relief

Acetaminophen 325 mg Pain reliever/Fever reducer Dextromethorphan HBr 10 mg Cough suppressant Phenylephrine HCl 5 mg Nasal decongestant

Relieves ✓ Aches, fever & sore throat ✓ Cough ✓ Nasal congestion

For ages 6 years and over Alcohol-free Antihistamine-free

## 12 fl oz (355 mL)

#### PEEL CORNER FOR MORE DRUG FACTS

#### Drug Facts (continued)

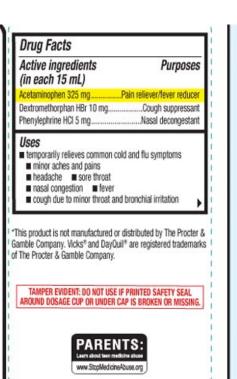
- If you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.
  Ask a doctor before use if you have
- liver disease heart disease
- high blood pressure thyroid disease
- diabetes a sodium-restricted diet
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with
- smoking, asthma, or emphysema cough that occurs with too much phlegm (mucus)

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin.

#### When using this product, do not exceed recommended dosage.

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
   pain, nasal congestion, or cough gets worse, or lasts more than 5 days (children) or 7 days (adults)
- 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.



#### Drug Facts (continued)

If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. Overdose warning: Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

PEEL CORNER FOR MORE DRUG FACTS

#### Directions

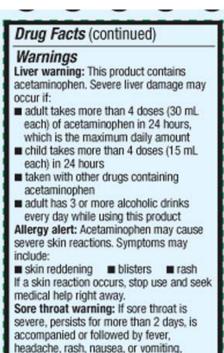
PLD-A380D LB009714

PLD-A380D

LB009717

- do not take more than directed (see Overdose warning)
- do not take more than 4 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other dosing
- device.
- mL = milliliter
- keep dosing cup with product

| adults and children<br>12 years and over                          | 30 mL every<br>4 hours                               |  |
|---|--|--|
| children 6 to<br>under 12 years<br>children 4 to<br>under 6 years | 15 mL every<br>4 hours<br>ask a doctor<br>do not use |  |
|   |  |  |



#### Drug Facts (continued)

consult a doctor promptly.

with any other drug containing

ask a doctor or pharmacist.

acetaminophen (prescription or

nonprescription). If you are not sure

whether a drug contains acetaminophen,

Do not use

Other information

each 15 mL contains: sodium 12 mg
 store between 20-25°C (68-77°F). Do not refrigerate.

**Inactive ingredients** citric acid, FD&C yellow #6, flavor, glycerin, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol, sucralose, xanthan gum

**QUALITY CHOICE Daytime Cold & Flu Relief** 

|  | -  | <b>FLU</b><br>rphan hbr phenylephrii | ae hel liquid  |                         |      |                    |  |
|--|--|--------------------------------------|----------------|-------------------------|------|--------------------|--|
| acetaminophen  | dextrometrio   |                                      |                |                         |      |                    |  |
| Product Info   | rmation  |                                      |                |                         |      |                    |  |
| Product Type   |  | HUMAN OTC DRUG                       | ltem Code      | em Code (Source)        |      | NDC:83324-167      |  |
| Route of Admir   | nistration   | ORAL                                 |                |                         |      |                    |  |
|  |  |                                      |                |                         |      |                    |  |
| Active Ingred  | lient/Active   | Moiety                               |                |                         |      |                    |  |
|  | Ingred   | dient Name                           |                | Basis of Stre           | ngth | Strength           |  |
| ACETAMINOPHEN  | <b>I</b> (UNII: 36209ITL   | .9D) (ACETAMINOPHEN - UNI            | ll:362O9ITL9D) | ACETAMINOPHEN           |      | 325 mg<br>in 15 mL |  |
| DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) DEXTROMETHORPHA<br>(DEXTROMETHORPHAN - UNII:7355X3ROTS) DEXTROMETHORPHA |  |                                      |                |                         | AN   | 10 mg<br>in 15 mL  |  |
|  | PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE -       PHENYLEPHRINE -         JNII:1WS297W6MV)       PHENYLEPHRINE - |                                      |                |                         |      | 5 mg<br>in 15 mL   |  |
| Inactive Indr  | edients  |                                      |                |                         |      |                    |  |
| Inactive Ingredients Ingredient Name   |  |                                      |                |                         |      | trength            |  |
| ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)   |  |                                      |                |                         |      |                    |  |
| FD&C YELLOW N  | I <b>O. 6</b> (UNII: H77V  | (EI93A8)                             |                |                         |      |                    |  |
| GLYCERIN (UNII: F  | DC6A3C0OX)   |                                      |                |                         |      |                    |  |
| PROPYLENE GLY  | COL (UNII: 6DC9  | Q167V3)                              |                |                         |      |                    |  |
| WATER (UNII: 059   | QF0KO0R)   |                                      |                |                         |      |                    |  |
| SACCHARIN SOD  | -  | ·                                    |                |                         |      |                    |  |
|  |  | E (UNII: B22547B95K)                 |                |                         |      |                    |  |
| SODIUM BENZOA  |  |                                      |                |                         |      |                    |  |
| SODIUM CHLORI  |  | /IQ8X)                               |                |                         |      |                    |  |
| SORBITOL (UNII: !  |  |                                      |                |                         |      |                    |  |
| SUCRALOSE (UNII<br>XANTHAN GUM (L  |  | =)                                   |                |                         |      |                    |  |
|  |  | -1                                   |                |                         |      |                    |  |
| Packaging  |  |                                      |                |                         |      |                    |  |
| # Item Code  |  | ackage Description                   |                | larketing Start<br>Date |      | eting End<br>Date  |  |
| <b>1</b> NDC:83324-<br>167-12  | 355 mL in 1 BO<br>Combination Pro  | TTLE, PLASTIC; Type 0: Not<br>oduct  | a 07           | /31/2024                |      |                    |  |
|  |  |                                      |                |                         |      |                    |  |
| Marketing  | Informat   | ion                                  |                |                         |      |                    |  |
| Marketing<br>Category  | Applica  | tion Number or Monog<br>Citation     | jraph M        | arketing Start<br>Date  |      | eting End<br>Date  |  |
| category   |  | Citation                             |                | Date                    |      | Date               |  |

Revised: 4/2024

QUALITY CHOICE (Chain Drug Marketing Association)