

**COLD AND FLU RELIEF DAYTIME- acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride liquid
Dolgenercorp, Inc. (DOLLAR GENERAL & REXALL)**

Drug Facts

Active ingredients (in each 15 mL)

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Phenylephrine 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
 - fever
 - nasal congestion
 - cough due to minor throat and bronchial irritation

Warnings

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

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

Allergy 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 the user has

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

Ask a doctor or pharmacist before use if the user is

taking the blood thinning drug warfarin.

When using this product,

do not exceed recommended dosage.

Stop use and ask a doctor if

- redness or swelling is present
- new symptoms occur
- nervousness, dizziness or sleeplessness occurs
- fever gets worse or lasts more than 3 days
- pain, nasal congestion or cough gets worse or lasts more than 5 days (children) or 7 days (adults)
- 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 Day Time and Night Time 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 **1-888-309-9030**

Principal Display Panel

Compare to the active ingredients in VICKS® DAYQUIL® Cold & Flu*

Daytime

Cold & Flu Relief

Acetaminophen 325 mg

Pain Reliever/Fever Reducer

Dextromethorphan HBr 10 mg

Cough Suppressant

Phenylephrine HCl 5 mg

Nasal Decongestant

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

For ages 6 years and over

- Alcohol free
- Antihistamine free
- Non-drowsy
- Multi-symptom

Original Flavor

FL OZ (mL)

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

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DISTRIBUTED BY OLD EAST MAIN CO.

100 MISSION RIDGE

GOODLETTSVILLE, TN 37072

Package Label

Drug Facts (continued)

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 (adult) ■ 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 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 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 other **Daytime** or **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 1-888-309-9030

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 100 MISSION RIDGE,
 GOODLETTSVILLE, TN 37072

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DG™ health

Compare to
 the active
 ingredients of
VICKS® DAYQUIL®
Cold & Flu*

Day Time Cold & Flu Relief

Acetaminophen 325 mg
Pain Reliever/Fever Reducer
Dextromethorphan HBr 10 mg
Cough Suppressant
Phenylephrine HCl 5 mg
Nasal Decongestant

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

For ages 6 years and over
 • Alcohol free • Antihistamine free
 • Non-drowsy • Multi-symptom

Original
 Flavor

12 FL OZ (355 mL)

Drug Facts

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

Active ingredients (in each 15 mL) **Purposes**
 Acetaminophen 325 mg.....Pain reliever/fever reducer
 Dextromethorphan HBr 10 mg.....Cough suppressant
 Phenylephrine HCl 5 mg.....Nasal decongestant

Uses ■ temporarily relieves common cold and flu symptoms
 ■ minor aches and pains ■ headache ■ fever ■ sore throat
 ■ nasal congestion ■ 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. **Allergy 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 your 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 ■ 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.

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.

DOLLAR GENERAL HEALTH Day Time Cold & Flu Relief

COLD AND FLU RELIEF DAYTIME

acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-366
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg in 15 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 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)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8Z UX40TY)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-366-12	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/31/2018	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/31/2018	

Labeler - Dolgencorp, Inc. (DOLLAR GENERAL & REXALL) (068331990)

Revised: 10/2023

Dolgencorp, Inc. (DOLLAR GENERAL & REXALL)