

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

Drug Facts

Active ingredients (in each 15 mL)

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Guaifenesin 200 mg

Phenylephrine HCl 5 mg

Purpose

Pain reliever/fever reducer

Cough suppressant

Expectorant

Nasal decongestant

Uses

- temporarily relieves common cold and flu symptoms:
 - nasal congestion
 - sinus congestion and pressure
 - minor aches and pains
 - headache
 - fever
 - sore throat
 - reduces swelling of nasal passages
 - cough due to minor throat and bronchial irritation
 - temporarily restores freer breathing through the nose
 - promotes nasal and or sinus drainage
 - helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make cough more productive

Warnings

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

- adult take 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
- blistere
- 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 is 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
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough as occurs with smoking, asthma, chronic bronchitis, or emphysema
- a sodium restricted diet

Ask a doctor or pharmacist before use if user is

taking the blood thinning drug warfarin

When using this product

do not take more than directed.

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)
- new symptoms occur
- fever gets worse or lasts more than 3 days
- redness or swelling is present

- 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 can cause serious 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 and for children even if you do not notice any signs or symptoms

Directions

- **take only as directed (see overdose warning)**
- do not take more than 4 doses in 24 hours
- measure only with dosing cup provided. Do not use any other dosing device
- mL = milliliter
- keep dosing cup with product
- **when using other Daytime or Nighttime products, carefully read each label to insure correct dosing**

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

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 of VICKS® DAYQUIL® Severe Cold & Flu Day Time

Severe Cold & Flu Relief

Acetaminophen 325 mg

Pain Reliever/Fever reducer

Dextromethorphan HBr 10 mg

Cough suppressant

Guaifenesin 200 mg

Expectorant

Phenylephrine HCl 5 mg

Nasal decongestant

- Headache, fever, sore throat, minor aches & pains
- Nasal / sinus congestion & sinus pressure
- Cough
- Chest congestion

For ages 6 years and over

- Max strength
- Alcohol free
- Non-drowsy

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

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

DISTRIBUTED BY OLD EAST MAIN CO.

100 MISSION RIDGE

GOODLETTSVILLE, TN 37072

Product Label

DG health

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

Day Time Severe Cold & Flu Relief



Acetaminophen 325 mg

Pain Reliever/Fever Reducer

Dextromethorphan HBr 10 mg

Cough Suppressant

Guaifenesin 200 mg

Expectorant

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Nasal Decongestant

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For ages 6 years and over

- Max strength
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8 FL OZ (237 mL)

PLD-A340B LB009969

Drug Facts

Active ingredients (in each 15 mL)

Acetaminophen 325 mg.....Pain reliever/fever reducer
 Dextromethorphan HBr 10 mg.....Cough suppressant
 Guaifenesin 200 mg.....Expectorant
 Phenylephrine HCl 5 mg.....Nasal decongestant

Uses

- temporarily relieves common cold/flu symptoms
- nasal congestion ■ sinus congestion and pressure
- minor aches and pains ■ headache ■ fever ▶

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PARENTS:

Learn about teen medicine abuse

www.StopMedicineAbuse.org

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Satisfaction
Guaranteed!
(888) 309-9030

PLD-A340B LB009969

PEEL CORNER FOR MORE DRUG FACTS ▶

Drug Facts (continued)

- sore throat
- reduces swelling of nasal passages
- cough due to minor throat and bronchial irritation
- temporarily restores freer breathing through the nose
- promotes nasal and/or sinus drainage
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

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- 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. ▶

PEEL CORNER FOR MORE DRUG FACTS ▶

Drug Facts (continued)

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

- 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 the user has

- liver disease ■ heart disease
- high blood pressure ■ thyroid disease
- diabetes ■ a sodium-restricted diet
- trouble urinating due to an enlarged prostate gland

Drug Facts (continued)

Ask a doctor or pharmacist before use if user is taking the blood thinning drug warfarin.

When using this product, do not take more than directed.

Stop use and ask a doctor if

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- new symptoms occur
- fever gets worse or lasts more than 3 days
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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 (1-800-368-1222) right

Drug Facts (continued)

Directions

- take only as 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
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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

■ cough that occurs with too much phlegm (mucus)
 ■ persistent or chronic cough as occurs with smoking, asthma, chronic bronchitis, or emphysema

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.

benzoate, sodium chloride, sodium citrate, sorbitol, sucralose, xanthan gum
Questions or comments?
 Call 1-888-309-9030

DOLLAR GENERAL HEALTH Daytime Severe Cold and Flu Relief

SEVERE COLD AND FLU RELIEF DAYTIME

acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-515
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
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 15 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-515-08	237 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: 9/2024

Dolgencorp, Inc. (DOLLAR GENERAL & REXALL)