

**SEVERE COLD AND COUGH RELIEF DAYTIME- acetaminophen,
dextromethorphan hbr, phenylephrine hcl liquid
Dolgenercorp, Inc. (DOLLAR GENERAL & REXALL)**

Drug Facts

Active ingredients (in each 30 mL)

Acetaminophen 650 mg

Dextromethorphan HBr 20 mg

Phenylephrine HCl 10 mg

Purposes

Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

Uses

- temporarily relieves these symptoms due to a cold
 - minor aches and pains
 - headache
 - nasal and sinus congestion
 - sore throat
 - cough due to minor throat and bronchial irritation
- temporarily reduces fever

Warnings

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

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 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 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
- diabetes
- high blood pressure
- thyroid disease
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema

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, cough, or nasal congestion 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 a 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 6 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 under 12 years of age: do not use

Other information

- **each 30 mL contains:** sodium 16 mg
- store between 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

acesulfame potassium, alcohol, citric acid, edetate disodium, FD&C blue #1, FD&C red #40, flavor, glycerin, maltitol, propylene glycol, purified water, sodium benzoate, sodium citrate

Principal Display Panel

Compare to active ingredients of Theraflu® ExpressMax® Daytime Severe Cold & Cough*

For Ages 12 Years & Over

DAYTIME

Severe Cold & Cough

Acetaminophen

Dextromethorphan HBr

Phenylephrine HCl

Pain Reliever / fever Reducer

Cough Suppressant

Nasal Decongestant

ALCOHOL 10%

FL OZ (mL)

Berry Flavor

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

*This product is not manufactured or distributed by GSK Consumer Healthcare, distributor of Theraflu® ExpressMax® Daytime Severe Cold & Cough.

DISTRIBUTED BY OLD EAST MAIN CO.

100 MISSION RIDGE
 GOODLETTSVILLE, TN 37072

Product Label

DG health

Compare to active ingredients of Theraflu® ExpressMax® Daytime Severe Cold & Cough*

For Ages 12 Years & Over

Day Time

Severe Cold & Cough

Acetaminophen

Dextromethorphan HBr
Phenylephrine HCl

Pain Reliever/Fever Reducer
Cough Suppressant
Nasal Decongestant

ALCOHOL 10%

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GOODLETTSVILLE, TN 37072

100* Satisfaction Guaranteed! (888) 309-9030



Berry Flavor

8.3 FL OZ (245 mL)

A1045 PLD-B398B LB008183



3 49580 50108 9

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PLD-B398B
LB004789

Drug Facts

| Active ingredients (in each 30 mL) | Purposes |
|--|-----------------------------|
| Acetaminophen 650 mg.....Pain reliever/fever reducer | Pain reliever/fever reducer |
| Dextromethorphan HBr 20 mg.....Cough suppressant | Cough suppressant |
| Phenylephrine HCl 10 mg.....Nasal decongestant | Nasal decongestant |

Uses

- temporarily relieves these symptoms due to a cold
- minor aches and pains ■ headache
- nasal and sinus congestion ■ sore throat
- cough due to minor throat and bronchial irritation
- temporarily reduces fever

PEEL CORNER FOR MORE DRUG FACTS

Drug Facts (continued)

Warnings

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

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 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 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

Drug Facts (continued)

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

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, cough, or nasal congestion gets worse or lasts more than 7 days
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- redness or swelling is present

Drug Facts (continued)

- 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 6 doses (180 mL) 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

Drug Facts (continued)

- adults and children 12 years and over
- 30 mL every 4 hours
- children under 12 years of age: do not use

Other information

- each 30 mL contains: sodium 16 mg
- store between 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

acesulfame potassium, alcohol, citric acid, disodium EDTA, FD&C blue #1, FD&C red #40, flavor, glycerin, maltitol, propylene glycol, purified water, sodium benzoate, sodium citrate

PEEL CORNER FOR MORE DRUG FACTS

DOLLAR GENERAL HEALTH Day Time Severe Cold & Cough Berry Flavor

SEVERE COLD AND COUGH RELIEF DAYTIME

acetaminophen, dextromethorphan hbr, phenylephrine hcl liquid

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:55910-801 |
| Route of Administration | ORAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|----------------------------------|--------------------|
| ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D) | ACETAMINOPHEN | 650 mg in 30 mL |
| DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 20 mg in 30 mL |
| PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV) | PHENYLEPHRINE HYDROCHLORIDE | 10 mg in 30 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| ACESULFAME POTASSIUM (UNII: 23OV73Q5G9) | |
| ALCOHOL (UNII: 3K9958V90M) | |
| ANHYDROUS CITRIC ACID (UNII: XF417D3PSL) | |
| EDETATE DISODIUM (UNII: 7FLD91C86K) | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |
| FD&C RED NO. 40 (UNII: WZB9127XOA) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| MALTITOL (UNII: D65DG142WK) | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | |
| WATER (UNII: 059QF0KO0R) | |
| SODIUM BENZOATE (UNII: OJ245FE5EU) | |
| TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K) | |

Product Characteristics

| | | | |
|-----------------|-------|---------------------|--|
| Color | | Score | |
| Shape | | Size | |
| Flavor | BERRY | Imprint Code | |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:55910-801-08 | 245 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 03/31/2017 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M012 | 03/31/2017 | |

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

Revised: 10/2023

Dolgencorp, Inc. (DOLLAR GENERAL & REXALL)