COLD TERMINATOR COLD RELIEF - phenylephrine hydrochloride, acetaminophen, dextromethorphan hydrobromide, guaifenesin tablet, film coated
Wildman Business Group

ACTIVE INGREDIENT

Active ingredients (in each caplet)

Acetaminophen 325 mg

Dextromethorphan Hydrobromide 10 mg

Guaifenesin 100 mg

Phenylephrine Hydrochloride 5 mg

PURPOSE

Purpose

Pain reliever/fever reducer

Cough suppressant

Expectorant

Nasal decongestant

INDICATIONS & USAGE

Uses

Temporarily relieves these symptoms due to the common cold:

- headache
- nasal congestion
- cough
- minor aches and pains
- sore throat
- sinus congestion and pressure
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes and make cough more productive

Temporarily reduces fever.

WARNINGS

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:
- blisters
- rash
- skin reddening

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

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 have ever had an allergic reaction to this product or any of its ingredients
- 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 DOCTOR

Ask a doctor before use if you have
- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- difficulty in urination due to enlargement of the prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough that lasts as occurs with smoking, asthma, chronic bronchitis, or emphysema

ASK DOCTOR/PHARMACIST

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

WHEN USING

When using this product do not exceed recommended dosage.

STOP USE

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 rash or headache that lasts.

These could be signs of a serious condition.

PREGNANCY OR BREAST FEEDING

If pregnant or breast-feeding, ask a health professional before use.

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

DOSAGE & ADMINISTRATION

Directions

- **do not take more than directed**

Adults and children: (12 years and over)

- take 2 caplets every 4 hours
- not to take more than 10 caplets in 24 hours

Children under 12 years: ask a doctor

OTHER SAFETY INFORMATION

Other information

- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- tamper evident sealed packets
- do not use any opened or torn packets

INACTIVE INGREDIENT

Inactive ingredients

corn starch, crospovidone, FD&C yellow #6 aluminum lake, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, talc, titanium dioxide

QUESTIONS

Questions or comments? (888) 602-0288

PRINCIPAL DISPLAY PANEL – 150 Tablet Box Label

Cold Terminator

Maximum Strength

Cold Relief

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Guaifenesin 100 mg

Phenylephrine HCl 5 mg

Pull to Open

RELIEVES

- Fever
- Pain
- Stuffy Nose
- Chest Congestion

The Provision

First Aid

Line™

BY WILDMAN

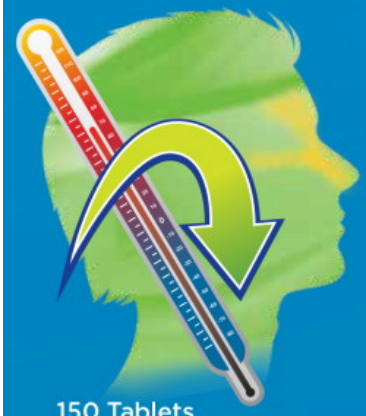
150 Caplets

2 per packet

2556

Cold Terminator
MAXIMUM STRENGTH
COLD RELIEF

Cold Terminator
MAXIMUM STRENGTH
COLD RELIEF



150 Tablets
2 Per Packet

RELIEVES:
- Fever
- Pain
- Stuffy Nose
- Chest Congestion

The Provision First Aid Line™
BY WILDMAN

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Guaifenesin 100 mg
Phenylephrine HCl 5 mg

RELIEVES:
- Fever
- Pain
- Stuffy Nose
- Chest Congestion

The Provision First Aid Line™
BY WILDMAN

150 Caplets

2 Per Packet

Manufactured for Provision by Wildman
Warsaw, IN 46580

8 58704 00423 3

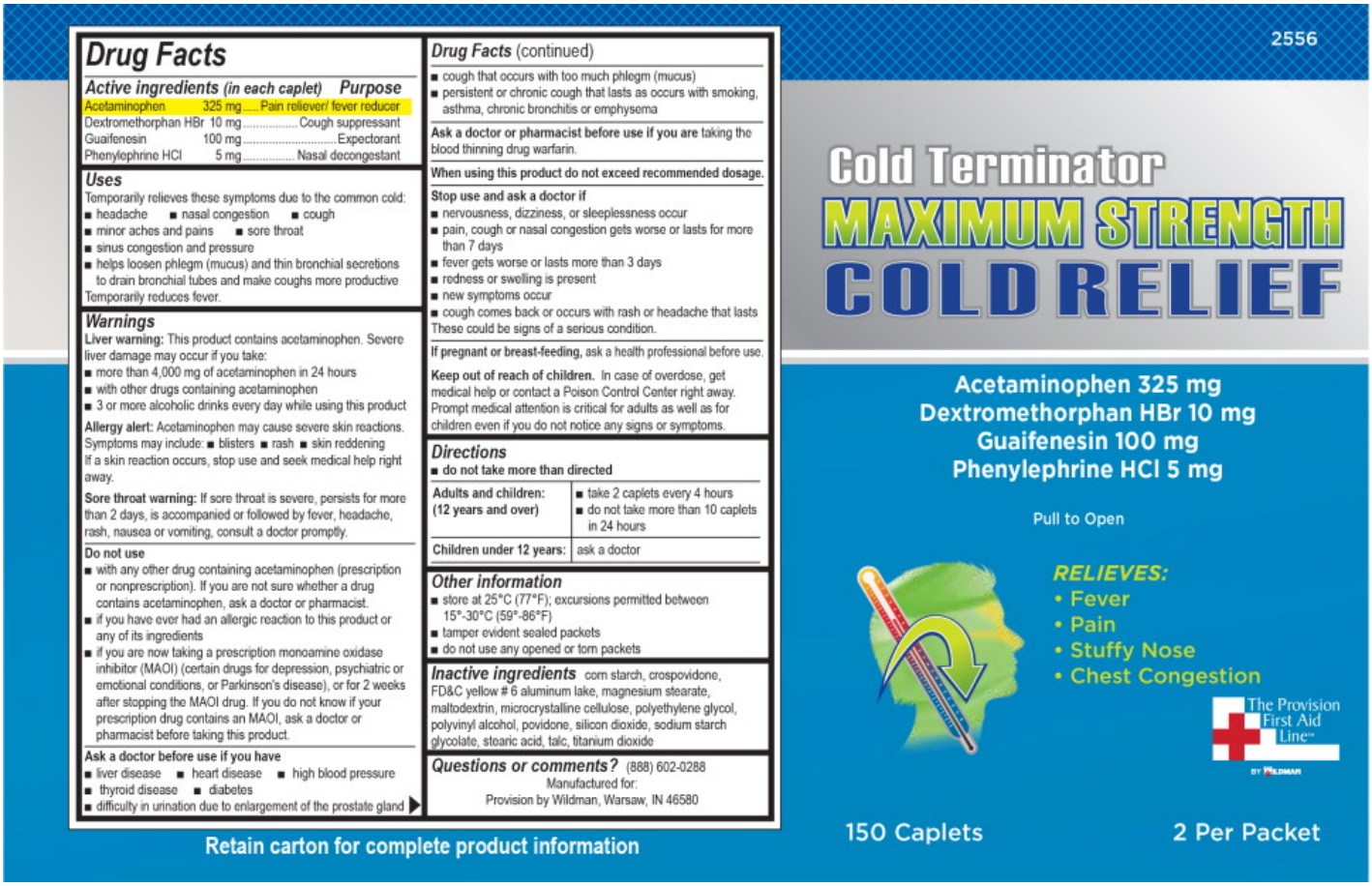
2556

Cold Terminator
MAXIMUM STRENGTH
COLD RELIEF

RELIEVES:
- Fever Reducer
- Pain
- Stuffy Nose
- Chest Congestion

The Provision First Aid Line™
BY WILDMAN

2556



COLD TERMINATOR COLD RELIEF

phenylephrine hydrochloride, acetaminophen, dextromethorphan hydrobromide, guaifenesin tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84269-2556
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CROSPOVIDONE (UNII: 2S7830E561)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

MALTODEXTRIN (UNII: 7CVR7L4A2D)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
POVIDONE (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	orange (orange)	**Score**	no score
Shape	CAPSULE (Caplet)	**Size**	17mm
Flavor		**Imprint Code**	44;546
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84269-2556-2	75 in 1 BOX	07/01/2024	
1		2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:84269-2556-1	50 in 1 BOX	07/01/2024	
2		2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/01/2024	

Labeler - Wildman Business Group (016677338)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	MANUFACTURE(84269-2556)

Establishment

Name	Address	ID/FEI	Business Operations
Medique Products		086911794	PACK(84269-2556)

Establishment

Name	Address	ID/FEI	Business Operations
Prestige Packaging		080667761	PACK(84269-2556)