PHYSICIANS CARE NON-DROWSY COLD AND COUGH- acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride tablet, film coated Acme United

Physicians Care Non-Drowsy Cold & Cough

Drug Facts

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Active Ingredient

Active ingredient (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

Uses

Uses

Temporarily relieves these cold symptoms

- cough
- sore throat
- minor aches and pains
- headache
- nasal congestion
- 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:

- 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

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 a doctor before use if you have

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

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• taking the blood thinning drug warfarin.

When using this product do not exceed recommended dosage.

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Stop use and ask a doctor if

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- new symptons occur
- redness or swelling is present
- pain, cough or nasal congestion gest worse or lasts for more than 7 days
- fever gets worse or lasts more than 3 days
- nervousness, dizziness, or sleeplessness 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.

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Keep out of reach of children.

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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.

Directions

Directions

do not take more than directed

Adults and children: (12 years and older)

Take 2 caplets every 4 hours

Do not to take more than 10 caplets in 24 hours

Children under 12 years: ask a doctor

Other 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 ingredients

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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? 1-800-835-2263

Physicians Care Cold & Cough

91462

Physicians Care®

Non-Drowsy

Cold & Cough

Cold, Cough and Flu Relief

Pull

to Open

COATED CAPLET

Individually packaged for workplace use

Tamper-Evident Packets of 2 Caplets

Do not use if packet is open or torn.

Not Actual Size

100 Caplets (50 Packets, 2 caplets each)



PHYSICIANS CARE NON-DROWSY COLD AND COUGH

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

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0924-0142	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg	
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg	

Inactive Ingredients			
Ingredient Name	Strength		
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)			
CROSPOVIDONE (UNII: 2S7830E561)			
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
STEARIC ACID (UNII: 4ELV7Z65AP)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)			
MALTODEXTRIN (UNII: 7CVR7L4A2D)			
POVIDONE (UNII: FZ989GH94E)			
STARCH, CORN (UNII: O8232NY3SJ)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)			
TALC (UNII: 7SEV7J4R1U)			

Product Characteristics			
Color	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:0924-0142- 01	50 in 1 BOX	01/19/2023	

	2 in 1 PACKET; Type 0: Not a Combination Product		
Marketing I	nformation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Managraph Drug	M012	01/19/2023	
OTC Monograph Drug	I MOTZ	01/19/2023	

Labeler - Acme United (001180207)

Registrant - Unifirst First Aid Corportaion (832947092)

Establishment			
Name	Address	ID/FEI	Business Operations
Prestige Packaging		080667761	pack(0924-0142)

Establishment			
Name	Address	ID/FEI	Business Operations
Medique Products		086911794	pack(0924-0142)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International		832867837	manufacture(0924-0142)

Revised: 5/2024 Acme United