#### HEAD CONGESTION PLUS FLU SEVERE PE- acetaminophen, dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride tablet, coated MEIJER DISTRIBUTION INC

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#### 1174-MEI-2022-0923

## **Drug Facts**

Active ingredients (in each caplet)	Purposes
Acetaminophen 325 mg	Pain reliever/fever reducer
Dextromethorphan HBr 10 mg	Cough suppressant
Guaifenesin 100 mg	Expectorant
Phenylephrine HCl 5 mg	Nasal decongestant

#### Uses

- temporarily relieves these symptoms due to the common cold:
  - nasal congestion
  - headache
  - minor aches and pains
  - cough
  - sore throat
  - sinus congestion and pressure
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes and make coughs more productive
- temporarily reduces fever

# Warnings

#### Liver warning

This product contains acetaminophen. The maximum daily dose of this product is 10 caplets (3,250 mg acetaminophen) in 24 hours. 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 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.
- if you have ever had an allergic reaction to this product or any of its ingredients

# 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
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, 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

#### Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- pain, nasal congestion, or cough 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.

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

#### Keep out of reach of children.

# **Overdose warning**

In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). 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)

adults and children 12 years and over	<ul> <li>take 2 caplets every 4 hours</li> <li>do not take more than 10 caplets in 24 hours</li> </ul>
children under 12 years	<ul> <li>ask a doctor</li> </ul>

#### Other information

- store between 20-25°C (68-77°F) in a dry place
- retain carton for complete product information and warnings

#### Inactive ingredients

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

#### **Questions or comments?**

1-844-705-4384

#### PRINCIPAL DISPLAY PANEL

meijer ®

NDC 41250-574-02

Compare to Sudafed PE® Head Congestion + Flu Severe active ingredients†

head congestion + flu severe PE

Acetaminophen | Dextromethorphan HBr | Guaifenesin | Phenylephrine HCl

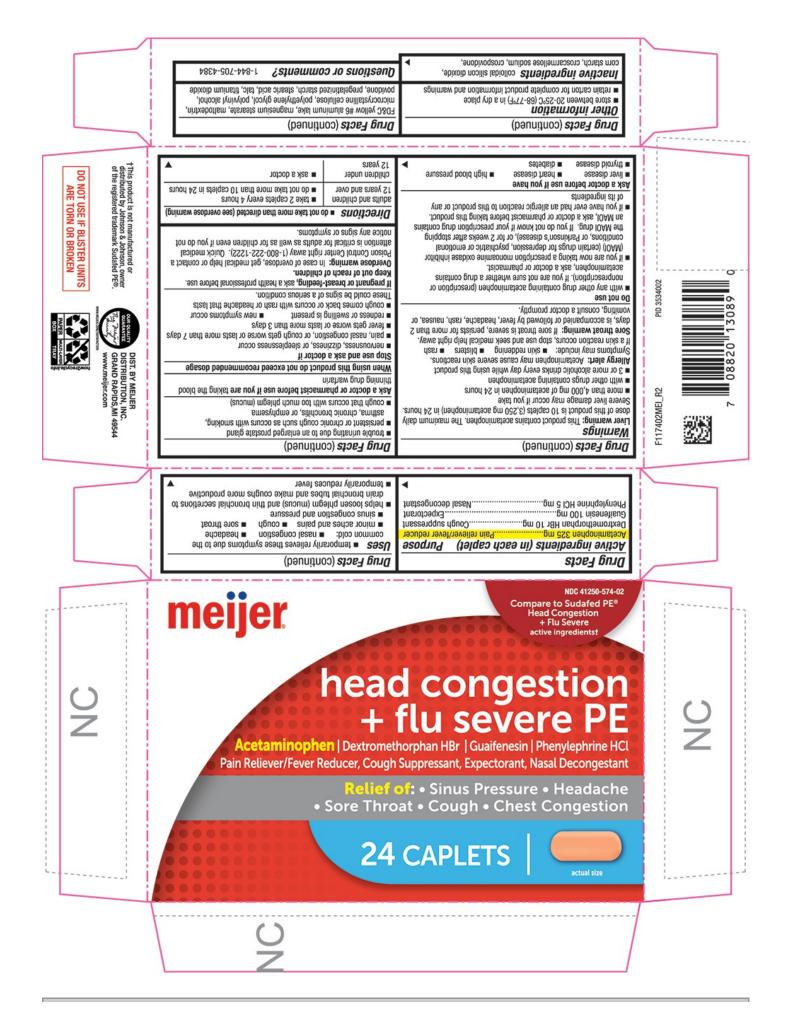
Pain Reliever/Fever Reducer, Cough Suppressant, Expectorant, Nasal Decongestant

Relief of: • Sinus Pressure • Headache

Sore Throat • Cough • Chest Congestion

24 CAPLETS

actual size



# HEAD CONGESTION PLUS FLU SEVERE PE

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

Product Information						
Product Type	HUMAN OT	C DRUG	Item Code (	Source)	NDC:4125	50-574
Route of Administration	ORAL					
Active Ingredient/Act	ive Moiety					
In	gredient Nan	ne		Basis of Str	ength	Strengt
ACETAMINOPHEN (UNII: 3620	D9ITL9D) (ACETA	MINOPHEN - UN	III:362O9ITL9D)	ACETAMINOPHEN		325 mg
DEXTROMETHORPHAN HYDI (DEXTROMETHORPHAN - UNII:7		NII: 9D2RTI9KYH	1)	DEXTROMETHORPH HYDROBROMIDE	HAN	10 mg
GUAIFENESIN (UNII: 495W745	1VQ) (GUAIFENE	SIN - UNII:495W	/7451VQ)	GUAIFENESIN		100 mg
PHENYLEPHRINE HYDROCHI UNII:1WS297W6MV)	LORIDE (UNII: 04	IJA59TNSJ) (PHE	NYLEPHRINE -	PHENYLEPHRINE HYDROCHLORIDE		5 mg
Inactive Ingredients						
	Ingred	lient Name			St	trength
SILICON DIOXIDE (UNII: ETJ72	Z6XBU4)					
CROSCARMELLOSE SODIUM	I (UNII: M28OL1H	H48)				
CROSPOVIDONE (UNII: 25783	30E561)					
FD&C YELLOW NO. 6 (UNII:	H77VEI93A8)					
ALUMINUM OXIDE (UNII: LMI2	2606933)					
MAGNESIUM STEARATE (UNI	I: 70097M6I30)					
MALTODEXTRIN (UNII: 7CVR7	L4A2D)					
CELLULOSE, MICROCRYSTA	LLINE (UNII: OPI	LR32D61U)				
POLYETHYLENE GLYCOL, U	NSPECIFIED (UN	NII: 3MJQ0SDW1	A)			
POLYVINYL ALCOHOL, UNSF	PECIFIED (UNII:	532B59J990)				
POVIDONE, UNSPECIFIED (L	JNII: FZ989GH94	E)				
STARCH, PREGELATINIZED (	CORN (UNII: 082	32NY3SJ)				
STEARIC ACID (UNII: 4ELV7Z	55AP)					
TALC (UNII: 7SEV7J4R1U)						
TITANIUM DIOXIDE (UNII: 15	FIX9V2JP)					
Product Characterist	ics					
Color	orange	Score		no	score	
Shape	OVAL	Size		17	'nm	
Flavor		Imprint C	Code	AA	A;1134	
Contains						
Packaging						
# Item Code	Package De	escription	Ма	rketing Start Date		ting End ate

<b>1</b> NDC:41250- 574-02 2	in 1 CARTON	09/23/2022	08/31/2026
	2 in 1 BLISTER PACK; Type 0: Not a Combination roduct		
Marketing I	nformation		
Marketing I Marketing Category	n <b>formation</b> Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Marketing	Application Number or Monograph Citation	-	-

Labeler - MEIJER DISTRIBUTION INC (006959555)

Revised: 10/2024

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