# DAYTIME SEVERE COLD AND FLU- acetaminophen, dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride tablet, film coated

**Spirit Pharmaceuticals LLC** 

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**Daytime Severe Cold and Flu** 

#### **Drug Facts**

## Active ingredients (in each caplet)

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Guaifenesin 200 mg
Phenylephrine HCl 5

#### **Purpose**

Pain reliever/fever reducer

Cough suppressant

Expectorant

Nasal decongestant

#### Uses

- for the temporary relief of the following cold/flu symptoms: minor aches and pains
- headache sore throat nasal congestion cough helps loosen phlegm (mucus) and thin

bronchial secretions to 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.

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

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adults and children 12 years & • take 2 caplets every 4 hours • swallow whole; do not crush, chew over children under 12 ask a doctor years
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#### Other information

store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

### **Inactive Ingredients**

croscarmellose sodium, D&C Yellow#10, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone\*, pregelatinized starch, talc, titanium dioxide \*may contain this ingredient

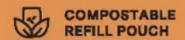
#### Questions or comments?

1-888-333-9792

## Distributed by:

Cabinet Health P.B.C.

## **Principal Display panel**





# CABINET:

# Daytime Severe Cold & Flu Refill

COMPARE TO THE ACTIVE INGREDIENTS IN:

Tylenol® Cold + Flu Severe\*

Acetaminophen 325 mg, Dextromethorphan HBr 10 mg, Gualfenesin 200 mg, Phenylephrine HCl 5 mg

#### RELIEVES

HEAD & BODY ACHES - FEVER & SORE THROAT - COUGH NASAL CONGESTION - MUCUS & CHEST CONGESTION





#### TO OPEN:

- 1. POSITION THUMBS INSIDE THE SHORT FLANGE OVER THE LOCKS.
- 2. GRIP WHILE PIVOTING THUMBS OUTWARD TO OPEN THE LOCKS.

Tamper evident: do not use if pouch is open

#### Drug Facts

#### Active Ingredient (in each caplet) Purpose .......... Pain reliever/fever reducer Acetaminophen 325 mg.......Pain reliever/fever reducer Dextromethorophan HBr 10 mg......Cough suppressant Gualfenesin 200 mg Expectorant Phenylephrine HCl 5 mg Nasal decongestant

**Uses** • for the temporary relief of the following cold/flu symptoms:
• minor aches and pains • headache • sore throat • nasal congestion . cough . helps loosen phiegm (mucus) and thin branchial secretions to 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 atcoholic drinks every day while using this product Allergy alert: acstaminophen may cause severe skin reactions. Symptoms may include: \* skin recidening \* bilsters \* 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 hed an allergic reaction to this product or any of its ingredients.

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

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#### **Drug Facts** (continued)

#### Directions a do not take more than directed

adults and children over 12 years old	take 2 caplets every 4 hours • swallow whole; do not crush, chew or dissolve     do not take more than 10 caplets in 24 hours
children under 12 years	ask a doctor

Other information • store at 25°C (77°F) excursions permitted between 15"-30°C (59"-86"F)

**Inactive Ingredients** Croscerneliose sodium, D&C Yellow#10, magnesium steerate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcehol, povidone\*, pregelatinized starch, talc, titanium dicode \*may contain this ingredient

#### Questions or comments? 1-898-333-9792

- This product is not manufactured or distributed by McNell Consumer Healthcare, distributor of Tylenol® Cold + Flu Severe
- Less than the limit of detection and consistent with gluten-free diet labeling per FDA



ACC-1-10033 Distributed by: Cabinet Health P.B.C. Question/Commente7 1-888-833-9792 www.cabinetheelth.com

Lot / Expiration w









## **DAYTIME SEVERE COLD AND FLU**

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

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68210-4221
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg	
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg	
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS 297W6MV)	PHENYLEPHRINE	5 mg	

Inactive Ingredients		
Ingredient Name	Strength	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)		
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)		
POVIDONE (UNII: FZ 989GH94E)		
STARCH, POTATO (UNII: 81089SAH3T)		
TALC (UNII: 7SEV7J4R1U)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		

Product Characteristics			
Color	yellow	Score	no score
Shape	CAPSULE	Size	20mm
Flavor		Imprint Code	ET32
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68210-4221- 3	30 in 1 JAR; Type 0: Not a Combination Product	11/14/2022	

## **Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	11/14/2022	

## Labeler - Spirit Pharmaceuticals LLC (179621011)

Revised: 12/2024 Spirit Pharmaceuticals LLC