# COLD AND FLU RELIEF DAYTIME- acetaminohpen, dextromethorphan hbr, phenylephrine hcl capsule, liquid filled Major Pharmaceuticals

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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### **Drug Facts**

# Active ingredients (in each softgel) Acetaminophen 325 mg

Dextromethoprhan HBr 10 mg
Phenlyephrine HCl 5 mg

## **Purpose**

## Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

#### Uses

- temporarily relieves common cold and flu symptoms:
  - minor aches and pains
  - headache
  - sore throat
  - nasal congestion
  - fever
  - cough due to minor throat and bronchial irritation

## **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 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
- diabetes
- heart disease
- thyroid disease
- high blood pressure
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, 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

- pain, cough, or nasal congestion gets worse or lasts more than 7 days
- nervousness, dizziness, or sleeplessness occur
- 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:** Taking more than the recommended dose can cause liver damage. In case of overdose, get medical help or contact a Poison Control Center 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 4 doses in 24 hours
- adults and children 12 years and over: take 2 softgels with water every 4 hours
- swallow whole; do not crush, chew, or dissolve
- children under 12 years: do not use
- when using other Daytime or Nighttime products, carefully read each label to insure correct dosing

#### Other information

- store at between 15-30°C (59-86°F)
- avoid excessive heat

## **Inactive ingredients**

butylated hydroxyanisole\*, butylated hydroxytoluene\*, FD&C red #40\*, FD&C yellow#6, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, sorbitan, sorbitol, titanium dioxide\*, white ink

\*contains one or more of these ingredients

### Questions or comments?

Call 1-877-753-3935 Monday-Friday 9AM-5PM EST

## **Principal Display Panel**

Compare to the active ingredients in Vick® DayQuil® Cold & Flu LiquiCaps®†

Daytime Multi-Symptom

Cold & Flu Relief

Acetaminophen 325 mg - Pain reliever/Fever reducer

Dextromethorphan HBr 10 mg Cough suppressant

Phenylephrine HCl 5 mg - Nasal decongestant

Non-Drowsy

Alcohol-free

Antihistamine-Free

Softgels\*\*

(\*\*Liquid-filled capsules)

†This product is not manufactured or distributed by the Procter & Gamble Company. Vicks®, DayQuil®, and LiquiCaps® are registered trademarks of The Procter & Gamble Company.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWING ANY SIGNS OF TAMPERING.

## KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Distributed by:

Major® Pharmaceuticals 17177 N laurel Park Drive, Suite 233 Livonia, MI 48152

## Package Label

"contains one or more of these ingredients

dioxide\*, white ink glycol, povidone, propylene glycol, purified water, sorbitan, sorbitol, titanium hydroxytoluene\*, FD&C red #40\*, FD&C yellow #6, gelatin, glycerin, polyethylene Inactive ingredients butylated hydroxyanisole\*, butylated

> ■ store between 15-30°C (59-86°F)
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**MAJOR**<sup>®</sup>

NDC 0904-6793-44

Compare to the active ingredients in Vicks® DayQuil® Cold & Flu LiquiCaps®†

Daytime Multi-Symptom

## Cold & Flu Relief

Acetaminophen 325 mg - Pain reliever/Fever reducer Dextromethorphan HBr 10 mg - Cough suppressant Phenylephrine HCl 5 mg - Nasal decongestant

Non-drowsy Alcohol-free Antihistamine-free

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Distributed by: Major® Pharmaceuticals

17177 N Laurel Park Drive, Suite 233 Livonia, MI 48152

Re-order No. 701001 R-58 Rev. 08/18 <sup>1</sup>This product is not manufactured or distributed by The Procter & Gamble Company, Vicks®, DayQuil®, and LiquiCaps® are registered trademarks of The Procter & Gamble Company.





#### **MAJOR Cold & Flu Relief**

## **COLD AND FLU RELIEF DAYTIME**

acetaminohpen, dextromethorphan hbr, phenylephrine hcl capsule, liquid filled

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0904-6793	

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	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg	

Inactive Ingredients		
Ingredient Name	Strength	
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)		
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)		
GELATIN (UNII: 2G86QN327L)		
GLYCERIN (UNII: PDC6A3C0OX)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POVIDONE (UNII: FZ 989GH94E)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SORBITAN (UNII: 6092ICV9RU)		
SORBITOL (UNII: 506T60A25R)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		

Product Characteristics				
Color	orange	Score	no score	
Shape	CAPSULE	Size	20mm	
Flavor		Imprint Code	P19;95A;512;AP016	

## Contains

F	Packaging				
#	tem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:0904-6793-	16 in 1 CARTON	08/31/2018	12/27/2024	
1	1	1 in 1 BLISTER PACK; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	08/31/2018	12/27/2024

## Labeler - Major Pharmaceuticals (191427277)

Revised: 9/2022 Major Pharmaceuticals