

**COLD AND FLU DAYTIME, NIGHTTIME, MULTI-SYMPTOM- acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl  
MEIJER DISTRIBUTION, INC**

*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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**Non-Drowsy Daytime & Nighttime Cold and Flu**

***Active ingredients (in each liquid-filled capsule) (Daytime Cold & Flu)***

Acetaminophen 325 mg  
Dextromethorphan HBr 10 mg  
Phenylephrine HCl 5 mg

***Purpose***

Pain reliever/fever reducer  
Cough suppressant  
Nasal decongestant

***Active ingredients (in each liquid-filled capsule) (Nighttime Cold & Flu)***

Acetaminophen 325 mg  
Dextromethorphan HBr 15 mg  
Doxylamine succinate 6.25 mg

***Purpose***

Pain reliever/fever reducer  
Cough suppressant  
Antihistamine

***Uses***

- temporarily relieves common cold and flu symptoms:
  - fever
  - headache
  - sore throat
  - minor aches and pains
  - cough due to minor throat and bronchial irritation
  - nasal congestion (***Daytime only***)
  - runny nose and sneezing (***Nighttime only***)

***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.
- if you have ever had an allergic reaction to this product or any of its ingredients
- to make a child sleepy (**Nighttime only**)

### **Ask a doctor before use if you have**

- cough that occurs with too much phlegm (mucus)
- liver disease
- difficulty in urination due to enlargement of the prostate gland
- heart disease (**Daytime only**)
- diabetes (**Daytime only**)
- thyroid disease (**Daytime only**)
- high blood pressure (**Daytime only**)
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema (**Daytime only**)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis, or emphysema (**Nighttime only**)
- glaucoma (**Nighttime only**)

### **Ask a doctor or pharmacist before use if you are**

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers (**Nighttime only**)

### **When using this product**

- **do not exceed recommended dosage**
- excitability may occur, especially in children (**Nighttime only**)

- avoid alcoholic beverages (**Nighttime only**)
- marked drowsiness may occur (**Nighttime only**)
- alcohol, sedatives, and tranquilizers may increase drowsiness (**Nighttime only**)
- use caution when driving a motor vehicle or operating machinery (**Nighttime only**)

### **Stop use and ask a doctor if**

- nervousness, dizziness, or sleeplessness occur (**Daytime only**)
- pain, nasal congestion, or cough gets worse or lasts more than 7 days (**Daytime only**)
- pain or cough gets worse or lasts more than 7 days (**Nighttime only**)
- 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 accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

### **If taking NIGHTTIME and DAYTIME products, carefully read each section to ensure correct dosing.**

#### **Directions (Daytime only)**

- **do not take more than directed**
- do not take more than 8 capsules per 24 hours
- adults and children 12 years and over: take 2 capsules with water every 4 hours
- children under 12 years: ask a doctor

#### **Directions (Nighttime only)**

- **do not take more than directed**
- do not take more than 8 capsules per 24 hours
- adults and children 12 years and over: take 2 capsules with water every 6 hours
- children under 12 years: ask a doctor

#### **Other information**

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°C-30°C (59°F-86°F)
- protect from heat, humidity and light
- see end flap for expiration date and lot number

***Inactive ingredients (Daytime only)***

FD&C Yellow No. 6, FD&C Red No. 40, gelatin, glycerin, polyethylene glycol 400, povidone, propylene glycol, purified water, sorbitol sorbitan, titanium dioxide

***Inactive ingredients (Nighttime only)***

D&C Yellow No. 10, FD&C Blue No. 1, gelatin, glycerin, polyethylene glycol 400, povidone, propylene glycol, purified water, sorbitol sorbitan, titanium dioxide

***Questions or comments?***

**1-888-333-9792**

**PRINCIPAL DISPLAY PANEL**

Compare to Vicks® DayQuil® &

NyQuil® Cold & Flu LiquiCaps®

active ingredients†

DAYTIME • NON DROWSY

Cold & Flu

ACETAMINOPHEN /

PAIN RELIEVER / FEVER REDUCER

DEXTROMETHORPHAN HBr /

COUGH SUPPRESSANT

PHENYLEPHRINE HCl /

NASAL DECONGESTANT

MULTI-SYMPTOM

ACTUAL

SIZE

32

LIQUID

CAPS

NIGHTTIME

Cold & Flu

ACETAMINOPHEN /

PAIN RELIEVER / FEVER REDUCER

DEXTROMETHORPHAN HBr /

COUGH SUPPRESSANT

DOXYLAMINE SUCCINATE /

ANTIHISTAMINE

MULTI-SYMP TOM

ACTUAL

SIZE

16

LIQUID

CAPS

48 TOTAL LIQUID CAPS

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED

OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY

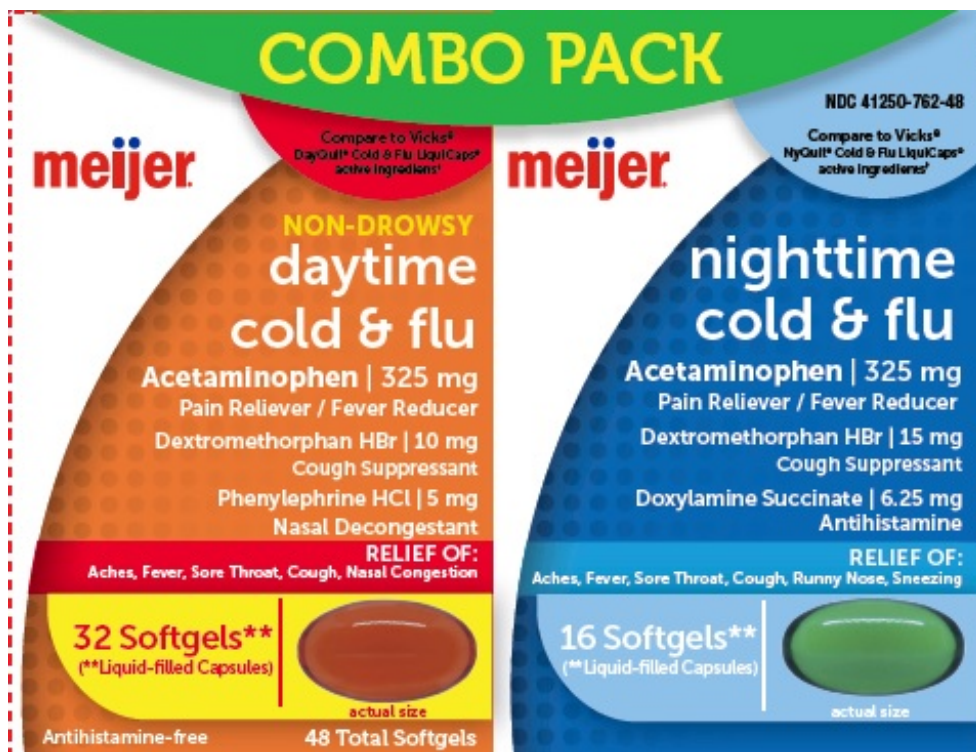
SIGNS OF TAMPERING

††This product is not manufactured or

distributed by Procter & Gamble, owner of the

registered trademark Vicks® DayQuil® &

NyQuil® Cold & Flu LiquiCaps®.



## COLD AND FLU DAYTIME, NIGHTTIME, MULTI-SYMP TOM

acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl kit

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:41250-762
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### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-762-48	1 in 1 PACKAGE, COMBINATION; Type 0: Not a Combination Product	09/16/2020	

### Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	2 BLISTER PACK	16
Part 2	4 BLISTER PACK	32

### Part 1 of 2

## COLD AND FLU NIGHTTIME, MULTI-SYMP TOM

acetaminophen, dextromethorphan hbr, doxylamine succinate capsule, liquid filled

### Product Information

Route of Administration ORAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg
<b>DOXYLAMINE SUCCINATE</b> (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>GELATIN</b> (UNII: 2G86QN327L)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>POVIDONE K30</b> (UNII: U725QWY32X)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>WATER</b> (UNII: 059QF0KO0R)	

### Product Characteristics

<b>Color</b>	green	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	21mm
<b>Flavor</b>		<b>Imprint Code</b>	215
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		8 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	09/16/2020	

Part 2 of 2

**COLD AND FLU DAYTIME, MULTI-SYMPATOM**

acetaminophen, dextromethorphan hbr, phenylephrine hcl capsule, liquid filled

## Product Information

**Route of Administration** ORAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

## Inactive Ingredients

Ingredient Name	Strength
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>POVIDONE K30</b> (UNII: U725QWY32X)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>GELATIN</b> (UNII: 2G86QN327L)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Product Characteristics

<b>Color</b>	red (ORANGE TO RED)	<b>Score</b>	no score
<b>Shape</b>	CAPSULE (Softgels)	<b>Size</b>	21mm
<b>Flavor</b>		<b>Imprint Code</b>	512
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		8 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	09/16/2020	



## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	09/16/2020	

**Labeler** - MEIJER DISTRIBUTION, INC (006959555)

Revised: 1/2023

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