

COLD AND FLU DAYTIME SEVERE- acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl tablet, film coated
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.

Meijer 44-640

Active ingredients (in each caplet)

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

Purpose

Pain reliever/fever reducer
Cough suppressant
Expectorant
Nasal decongestant

Uses

- temporarily relieves common cold and flu symptoms:
 - minor aches and pains
 - sinus congestion and pressure
 - sore throat
 - fever
 - headache
 - nasal congestion
 - cough due to minor throat and bronchial irritation
- reduces swelling of nasal passages
- temporarily restores freer breathing through the nose
- promotes nasal and/or sinus drainage
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

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:

- blisters
- rash
- skin reddening

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
- thyroid disease
- diabetes
- heart disease
- high blood pressure
- difficulty in urination due to enlargement of the prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough such 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.

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 accidental 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

- **do not take more than directed**
- do not take more than 8 caplets in 24 hours
- adults and children 12 years and over: take 2 caplets with water every 4 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°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

corn starch, crospovidone, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polysorbate 80, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, talc, titanium dioxide

Questions or comments?

1-800-426-9391

Principal Display Panel

meijer®

NDC 41250-643-08

**Compare to
Vicks® DayQuil®
Severe Cold & Flu
active ingredients***

**NON-DROWSY
MAXIMUM STRENGTH
daytime
severe cold & flu**

Acetaminophen | Pain Reliever | Fever Reducer
Dextromethorphan HBr | Cough Suppressant
Guaifenesin | Expectorant
Phenylephrine HCl | Nasal Decongestant

Relieves: Headache, Fever, Sore Throat, Minor Aches & Pains,
Nasal/Sinus Congestion & Sinus Pressure, Cough, Chest Congestion

24 Caplets

Actual
Size

**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 The
Procter & Gamble Company, owner of the registered
trademark Vicks® DayQuil® Severe Cold & Flu.
50844 ORG051964008

**DIST. BY MEIJER
DISTRIBUTION, INC.
GRAND RAPIDS, MI 49544
www.meijer.com**

meijer

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MAXIMUM STRENGTH**

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Drug Facts

**KEEP OUTER PACKAGE FOR
COMPLETE PRODUCT INFORMATION**

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Drug Facts (continued)

Uses ■ temporarily relieves common cold and flu symptoms:
■ minor aches and pains ■ sinus congestion and pressure
■ sore throat ■ headache ■ fever
■ cough due to minor throat and bronchial irritation
■ reduces swelling of nasal passages
■ temporarily restores free breathing through the nose
■ promotes nasal and/or sinus drainage

Drug Facts (continued)

■ helps loosen phlegm (mucus) and thin bronchial secretions to rid the
productive
bronchial passageways of bothersome mucus and make coughs more

Warnings

Liver warning: This product contains acetaminophen. Severe liver
damage may occur if you take
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■ 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: ■ rash ■ blisters ■ skin redness
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

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nonprescription). If you are not sure whether a drug contains
acetaminophen, ask a doctor or pharmacist.
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(MAOI) (certain drugs for depression, psychiatric or emotional
conditions, or Parkinson's disease), or for 2 weeks after stopping the
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ingredients
Ask a doctor before use if you have
■ liver disease ■ thyroid disease ■ diabetes
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■ difficulty in urination due to enlargement of the prostate gland

Drug Facts (continued)

Inactive ingredients
corn starch, croscopolone, FD&C red #40
aluminum lake, FD&C yellow #6 aluminum lake, magnesium stearate,
maltodextrin, microcrystalline cellulose, polyethylene glycol, polyorbate
80, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate,
stearic acid, talc, titanium dioxide

(Questions or comments?) 1-800-426-9391

**DIST. BY MEIJER
DISTRIBUTION, INC.
GRAND RAPIDS, MI 49544**



50844 ORG051964008
Procter & Gamble Company, owner of the registered
trademark Vicks® DayQuil® Severe Cold & Flu.
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Other information

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BLISTER IS TORN OR BROKEN
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4 hours
■ children under 12 years: ask a doctor
do not notice any signs or symptoms.

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 right away. Prompt
medical attention is critical for adults as well as for children even if you

could be signs of a serious condition.
■ cough comes back or occurs with rash or headache that lasts. These
■ redness or swelling is present ■ new symptoms occur
■ fever gets worse or lasts more than 3 days
■ pain, nasal congestion, or cough gets worse or lasts more than 7 days
■ nervousness, dizziness, or sleeplessness occur

Stop use and ask a doctor if
When using this product do not exceed recommended dosage.

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

Drug Facts (continued)

■ cough that occurs with too much phlegm (mucus)
■ persistent or chronic cough such as occurs with smoking, asthma,
chronic bronchitis, or emphysema

**TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER
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**No Print/No Varnish
Lot and Expiration No.**

PID 3416663



8-1214-640-08
ORG051964008

COLD AND FLU DAYTIME SEVERE

acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-643
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CROSPVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	ORANGE	Score	no score
Shape	OVAL	Size	19mm
Flavor		Imprint Code	44;640
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-643-08	2 in 1 CARTON	05/08/2020	
1		12 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	05/08/2020	

Labeler - Meijer Distribution Inc (006959555)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	MANUFACTURE(41250-643)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	PACK(41250-643)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	PACK(41250-643)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	PACK(41250-643)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	PACK(41250-643)

Revised: 3/2020

Meijer Distribution Inc