

DAYTIME COLD AND FLU- acetaminophen, dextromethorphan, phenylephrine capsule, liquid filled

Mckesson

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.

Acetaminophen, Dextromethorphan and Phenylephrine Day time Cold and Flu

Active Ingredient

(in each softgel)

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Phenylephrine HCl 5 mg

Purpose

pain reliever

Cough Suppressant

Nasal decongestant

Uses

pain reliever, cough suppressant and Nasal decongestant

Warnings

Warnings Failure to follow these warnings could result in serious consequences.

Liver Warning: This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4 doses in 24 hours which is maximum daily amount for this product
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks daily while using this product

do not use:

- with any other drug containing acetaminophen (prescription or not prescription). 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
- Heart disease
- Thyroid disease
- Diabetes
- High blood pressure
- Trouble urinating due to enlarged prostate gland

ask your doctor or pharmacist before use if you are taking the blood thinning drug warfarin.

When using this product, do not use more than directed.

Stop use and ask a doctor if:

- Redness or swelling is present
- You get nervous, dizzy or sleepless
- Fever gets worse or lasts more than 3 days
- New symptoms occur
- Symptoms do not get better within 7 days or are accompanied by a fever

Keep out of reach of children.

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

Overdose warning: Taking more than recommended dose (overdose) may cause liver damage. In case of 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.

Tamper evident: this package is safety sealed and child resistant. Use only if blisters are intact. If difficult to open use scissors.

Direction

- **do not exceed 4 doses per 24 hours**
- **take only as directed – see overdose warning**
- **Adults and children 12 years and over:** 2 softgels with water every 4 hours
- **Children under 4 to under 12 years:** ask a doctor
- **Children under 4 years:** do not use

Other Information

- store at room temperature

Inactive Ingredients

FD&C Red No.40, FD&C Yellow No. 6, Gelatin, Glycerin, Polyethylene Glycol, Povidone, Propylene Glycol, Purified Water, Sorbitol Special, Titanium dioxide

Questions or Comments

Call toll free 1-855-314-1850

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

Another Quality Product Distributed By McKesson
 One Post Street, San Francisco, CA 94104
 Money Back Guarantee
 Please visit us at www.sunmarkbrand.com

McKesson



Made in India

24 SOFTGELS (liquid-filled capsules)
COLD & FLU MULTI-SYMPOM



Non-drowsy
day time liquid caps
 Pain reliever, Fever reducer,
 Cough suppressant,
 Nasal decongestant,
 Alcohol-free
 Antihistamine-free
 DEXTROMETHORPHAN HBr,
 PHENYLEPHRINE HCl

Non-drowsy

day time liquid caps

sunmark®

COMPARE TO "VICKS® DAYQUIL®"
 ACTIVE INGREDIENT™
 NDC 70677-0011-1

sunmark®
 day time liquid caps

Drug Facts

Active ingredients (in each softgel) Purpose
 Acetaminophen 325 mg..... Pain reliever/Fever reducer
 Dextromethorphan HBr 10 mg..... Cough suppressant
 Phenylephrine HCl 5 mg..... Nasal decongestant

Directions

■ Do not exceed 4 doses per 24 hours.
 ■ Take only as directed - see Overdose warning
 Adults and Children 12 years and over ■ 2 softgels with water every 4 hours
 Children 4 to under 12 years ■ ask a doctor
 Children under 4 years ■ do not use

Inactive ingredients: FD&C Re No. 40, FD&C Yellow No. 6, Gelatin, Glycerin, Polyethylene Glycol, Povidone, Propylene Glycol, Purified Water, Sorbitol, Saccharin, Titanium Dioxide

Other Information: Store at room temperature

Questions? call 1-855-314-1850

Drug Facts (continued)

Uses: temporarily relieves common cold/flu symptoms:
 ■ nasal congestion ■ cough due to minor throat & bronchial irritation ■ sore throat ■ headache ■ minor aches & pains ■ fever

Warnings: Failure to follow these warnings could result in serious consequences.
Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take: ■ More than 4 doses in 24 hours, which is the maximum daily amount for this product ■ With other drugs containing acetaminophen ■ 3 or more alcoholic drinks daily while using this product

Allergy alert: acetaminophen may cause severe skin reactions, symptoms may include: ■ skin redness ■ 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 ■ Heart disease ■ Thyroid disease ■ Diabetes ■ High blood pressure
 ■ Trouble urinating due to enlarged prostate gland ■ cough that occurs with too much phlegm (mucus)
 ■ Persistent or chronic cough such as occurs with smoking, asthma, or emphysema

Ask a doctor or pharmacist before use if you are taking the blood-thinning drug warfarin.
When using this product, do not use more than directed.
Stop use and ask a doctor if: ■ Redness or swelling is present ■ You get nervous, dizzy, or sleepless ■ New symptoms occur
 ■ Fever gets worse or lasts more than 3 days ■ pain, nasal congestion or cough get worse or last more than 7 days for an adult or 5 days for a child 12 years or older ■ 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 serious health problems. In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical for adults and for children even if you do not notice any signs or symptoms.

NDC: 70677-0011-1 24 liquid caps

DAYTIME COLD AND FLU

acetaminophen,dextromethorphan,phenylephrine capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70677-0011
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
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
POLYETHYLENE GLYCOL 1000 (UNII: U076Q6Q621)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITOL (UNII: 506T60A25R)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	ORANGE	Score	no score
Shape	CAPSULE	Size	22mm
Flavor		Imprint Code	512
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70677-0011-1	24 in 1 BLISTER PACK	11/25/2016	
1		1 in 1 CARTON; 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	11/25/2016	

Labeler - Mckesson (177667227)

Registrant - Velocity Pharma (962198409)

Revised: 12/2016

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