NIGHTIME COLD AND FLU- acetaminophen, dextromethorphan hydrobromide, doxylamine succinate capsule, liquid filled
Meijer, Inc.

Nighttime Cold and Flu

Drug Facts

Active ingredients (in each softgel)

Acetaminophen 325 mg

Dextromethorphan hydrobromide 15 mg

Doxylamine succinate 6.25 mg

Purpose

Pain reliever/fever reducer

Cough suppressant

Antihistamine

Uses

temporarily relieves common cold/flu symptoms:
• cough due to minor throat and bronchial irritation
• sore throat n headache n minor aches and pains
• fever
• runny nose and sneezing

Warnings

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 reddening
• blisters
• rash
If a skin reaction occurs, stop use and seek medical help right away.

Sore throat warning: If sore throat is severe, lasts for more than 2 days, or occurs with or is followed by fever, headache, rash, nausea, or vomiting, see 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
• glaucoma
• cough that occurs with too much phlegm (mucus)
• a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis, or emphysema
• trouble urinating due to enlarged prostate gland

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

When using this product
• do not use more than directed
• excitability may occur, especially in children
• marked drowsiness may occur n avoid alcoholic drinks
• be careful when driving a motor vehicle or operating machinery
• alcohol, sedatives and tranquilizers may increase drowsiness

Stop use and ask a doctor if
• pain 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.

Overdose warning: Taking more than directed 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.

Directions
- take only as directed - **see Overdose warning.**
- do not exceed 4 doses per 24 hours

adults & children 12 years & over	2 softgels with water every 6 hours
children 4 to under 12 years	ask a doctor
children under 4 years	do not use

When using other Daytime or Nighttime products, carefully read each label to ensure correct dosing

Other information

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

Inactive Ingredients

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

Questions or comments?

1-888-333-9792

Principal Display Carton

NIGHTIME COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79481-0027
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	15 mg
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITOL (UNII: 506T60A25R)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	

Product Characteristics

Color	green	**Score**	no score
Shape	OVAL	**Size**	21mm
Flavor		**Imprint Code**	902;215;AP02
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79481-0027-2	2 in 1 CARTON	09/15/2022	
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 Drug	M012	09/15/2022	

Labeler - Meijer, Inc. (006959555)

Revised: 12/2023 Meijer, Inc.