VAPOR ICE DAYTIME AND NITETIME COLD AND FLU- acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride,guaifenesin Meijer, 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 Distribution, Inc. Vapor Ice[™] Daytime & Nitetime Cold & Flu Drug Facts

Active ingredients (in each 30 mL) - Nitetime

Acetaminophen 650 mg

Dextromethorphan HBr 20 mg

Doxylamine succinate 12.5 mg

Phenylephrine HCl 10 mg

Purpose

Pain reliever/fever reducer

Cough suppressant

Antihistamine

Nasal decongestant

Uses

temporarily relieves common cold/flu symptoms:

- nasal congestion
- sinus congestion and pressure
- minor aches and pains
- headache
- fever
- sore throat
- runny nose and sneezing
- cough due to minor throat and bronchial irritation
- cough to help you sleep
- reduces swelling of nasal passages
- promotes nasal and/or sinus drainage
- temporarily restores freer breathing through the nose

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

Ask a doctor before use if you have

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- glaucoma
- cough that occurs with too much phlegm (mucus)
- a breathing problem such as emphysema or chronic bronchitis
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema
- sodium-restricted diet

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
- 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

- you get nervous, dizzy or sleepless
- 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.

Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

- take only as directed see Overdose warning
- only use the dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over	30 mL every 4 hrs
children 4 to under 12 yrs	ask a doctor
children under 4 yrs	do not use

Other information

- each 30 mL contains: sodium 42 mg
- store at 20-25°C (68-77°F)

Inactive ingredients

alcohol, anhydrous citric acid, D&C yellow #10, edetate disodium, FD&C blue #1, flavor, glycerin, polyethylene glycol, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol solution, sucralose

Questions or comments? 1-800-719-9260

Active ingredients (in each 30 mL) - Daytime

Acetaminophen 650 mg Dextromethorphan HBr 20 mg Guaifenesin 400 mg Phenylephrine HCl 10 mg

Purpose

Pain reliever/fever reducer

Cough suppressant

Expectorant

Nasal decongestant

Uses

temporarily relieves common cold/flu symptoms:

- nasal congestion
- sinus congestion and pressure
- cough due to minor throat and bronchial irritation
- minor aches and pains
- headache
- fever
- sore throat
- 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

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- with other drugs containing acetaminophen
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Do not use

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- 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
- heart disease
- thyroid disease
- diabetes
- high blood pressure
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- a sodium-restricted diet

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

- you get nervous, dizzy or sleepless
- 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.

Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

- take only as directed see Overdose warning
- only use the dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over	30 mL every 4 hrs
children 4 to under 12 yrs	ask a doctor
children under 4 yrs	do not use

Other information

- each 30 mL contains: sodium 41 mg
- store at 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

alcohol, anhydrous citric acid, D&C yellow #10, edetate disodium, FD&C blue #1, flavor, glycerin, polyethylene glycol, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol solution, sucralose

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

DAYTIME & NITETIME COMBO PACK

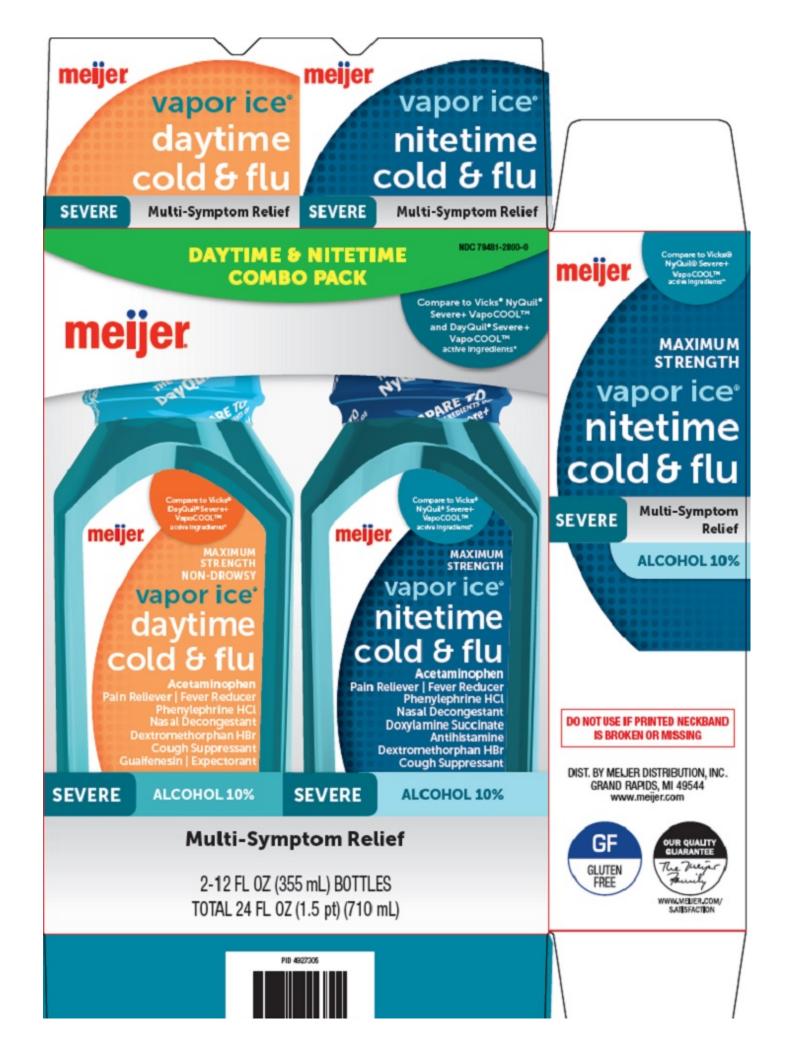
Compare to Vicks[®] NyQuil[®] Severe+VapoCOOL[™] and DayQuil[®] Severe+VapoCOOL[™]

active ingredients

meijer®

Compare to Vicks[®] DayQuil[®] Severe+VapoCOOL[™] active ingredients

meijer_® MAXIMUM STRENGTH **NON-DROWSY** vapor ice[®] daytime cold & flu Acetaminophen Pain Reliever | Fever Reducer Phenylephrine HCl Nasal Decongestant Dextromethorphan HBr **Cough Suppressant** Guaifenesin | Expectorant SEVERE ALCOHOL 10% Compare to Vicks[®] NyQuil[®] Severe+VapoCOOL[™] active ingredients meijer_® MAXIMUM STRENGTH vapor ice[®] nitetime cold & flu Acetaminophen Pain Reliever | Fever Reducer Phenylephrine HCl Nasal Decongestant **Doxylamine Succinate** Antihistamine Dextromethorphan HBr **Cough Suppressant SEVERE** ALCOHOL 10% Multi-Symptom Relief 2 -12 FL OZ (355 mL) BOTTLES TOTAL 24 FL OZ (1.5 pt) (710 mL)





28R02 6E C2



Vapor Ice® Niletime Severe	Vapor Ice® Daytime Severe	_
Drug Facts	Drug Facts	
Active ingredients (in each 30 mL) Purpose		moji
Acetaminophen (50 mgPain relevenfever reducer	Acetaminophen 850 mgPainrelieven/lever reduc	
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3 or more alsoholis drinks every day while using this product Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:	Il more than 4,000 mg of acetaminophen in 24 hours Il with other drugs containing acetaminophen	
Eskin reddening Ebisters Erash	■ 3 or more alcoholic drinks every day while using this product	
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	Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include: It skin reddening III blisters III nah	10000
or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.	If a skin reaction occurs, stop use and seek medical help right away.	
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E redness or swelling is present E new symptoms occur E cough comes back or occurs with rach or headache that lasts.	These sould be signs of a serious condition.	www.S
These could be signs of a serious condition.	If pregnant or breast-feeding, ask a health professional before use.	
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meijer MAXIMUM STRENGTH NON-DROWSY Vapor ice daytime cold & flu

Multi-Symptom Relief

ALCOHOL 10%





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VAPOR ICE DAYTIME AND NITETIME COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride,guaifenesin kit

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79481-2800

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:79481- 2800-0	1 in 1 PACKAGE; Type 0: Not a Combination Product	09/23/2021	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	355 mL
Part 2	1 BOTTLE	355 mL

Part 1 of 2

VAPOR ICE NITETIME COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride solution

Product Information	
Item Code (Source)	NDC:79481-0590
Route of Administration	ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	650 mg in 30 mL	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 30 mL	
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 30 mL	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS 297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 30 mL	

Inactive Incure				
Inactive Ingre	dients			
		Ingredient Name		Strength
ALCOHOL (UNII: 3k				
ANHYDROUS CITR				
D&C YELLOW NO				
EDETATE DISODIU				
D&C BLUE NO. 1		BD)		
GLYCERIN (UNII: PI				
		CIFIED (UNII: 3WJQ0SDW1A)		
		167V3)		
WATER (UNII: 0590		N/AOTN)		
SACCHARIN SODI				
		FORM (UNII: 1Q73Q2JULR)		
SORBITOL (UNII: 5				
SUCRALOSE (UNII:	96K6UQ3ZD4)			
Packaging				
# Item Code	Pac	kage Description	Marketing Start Date	Marketing End Date
1 NDC:79481- 0590-0	355 mL in 1 BO ⁻ Product	TTLE; Type 0: Not a Combination		
		on		
Marketing	informati			
Marketing Marketing			Marketing Start	Marketing End
•		ion Number or Monograph Citation	Marketing Start Date	Marketing End Date
Marketing Category	Applicat	ion Number or Monograph	-	-
Marketing Category	Applicat	ion Number or Monograph	-	-
Marketing Category OTC monograph fin	Applicat	ion Number or Monograph	-	-
	Applicat	ion Number or Monograph	-	-
Marketing Category OTC monograph fin Part 2 of 2	Applicat	ion Number or Monograph Citation	-	-
Marketing Category OTC monograph fin Part 2 of 2 VAPOR ICE	Application of the second seco	ion Number or Monograph Citation	Date	Date
Marketing Category DTC monograph fin Part 2 of 2 VAPOR ICE acetaminophen	Application of the second seco	ion Number or Monograph Citation	Date	Date
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Marketing Category DTC monograph fin Part 2 of 2 VAPOR ICE acetaminophen	Application of the second seco	ion Number or Monograph Citation	Date	Date
Marketing Category DTC monograph fin Part 2 of 2 VAPOR ICE acetaminophen solution	Application al part341 DAYTIME , dextrometho	ion Number or Monograph Citation	Date	Date
Marketing Category OTC monograph fin Part 2 of 2 VAPOR ICE acetaminophen solution	Application	ion Number or Monograph Citation	Date	Date
Marketing Category OTC monograph fin Part 2 of 2 VAPOR ICE	Application part341	ion Number or Monograph Citation	Date	Date

Active Ingredient/A	Active Moiety
	Ingredient Name

Basis of Strength Strength

ACETAMINOPHEN	(UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209	ITL9D) ACETAMINOPHEN	650 mg in 30 mL
	PHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) IAN - UNII:7355X3ROTS)	DEXTROMETHORPH HYDROBROMIDE	AN 20 mg in 30 mL
GUAIFENESIN (UN	II: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 30 mL
PHENYLEPHRINE UNII:1WS297W6MV)	HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHF	RINE - PHENYLEPHRINE HYDROCHLORIDE	10 mg in 30 mL
Inactive Ingre	edients		
	Ingredient Name		Strength
ALCOHOL (UNII: 3k	(9958V90M)		
ANHYDROUS CITR	RIC ACID (UNII: XF417D3PSL)		
D&C YELLOW NO	. 10 (UNII: 35SW5USQ3G)		
EDETATE DISODIU	JM (UNII: 7FLD91C86K)		
FD&C BLUE NO. 1	L (UNII: H3R47K3TBD)		
GLYCERIN (UNII: PI			
	GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
PROPYLENE GLYC	COL (UNII: 6DC9Q167V3)		
WATER (UNII: 0590	QF0KO0R)		
	UM (UNII: SB8ZUX40TY)		
	TE (UNII: OJ245FE5EU)		
	DE (UNII: 451W47IQ8X)		
	, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)		
SORBITOL (UNII: 5			
SUCRALOSE (UNII:	96K6UQ3ZD4)		
Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:79481- 0940-0	355 mL in 1 BOTTLE; Type 0: Not a Combination Product		
Marketing	Information		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph fir	nal part341		
Marketing	Information		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph fir	nal part341	09/23/2021	

Revised: 11/2022