### SEVERE NIGHTTIME SEVERE DAYTIME- acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl CVS Pharmacy

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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### CVS Pharmacy, Inc. Daytime Nighttime Cold & Flu Relief Drug Facts

## Nighttime Severe Cold & Flu Active ingredients (in each 30 mL)

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
- a 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 41 mg
- store at 20-25°C (68-77°F)

#### **Inactive ingredients**

anhydrous citric acid, edetate disodium, FD&C blue #1, FD&C red #40, flavor, glycerin, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol solution, sucralose, xanthan gum

#### Questions or comments?

1-800-719-9260

### Daytime Severe Cold & Flu Active ingredients (in each 15 mL)

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/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**

Liver warning: This product contains acetaminophen. Severe liver damage may occur if

- adult takes more than 4,000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours
- taken with other drugs containing acetaminophen
- adult has 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
- 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

#### 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 5 days (children) or 7 days (adults)
- 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 care 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 6 to under 12 yrs	15 mL every 4 hrs
children 4 to under 6 yrs	ask a doctor
children under 4 yrs	do not use

#### Other information

- each 15 mL contains: sodium 6 mg
- store at 20-25°C (68-77°F). Do not refrigerate.

#### **Inactive ingredients**

butylated hydroxyanisole, edetate disodium, FD&C yellow #6, flavor, glycerin, menthol, monobasic sodium phosphate, polyethylene glycol, propylene glycol, purified water, saccharin sodium, sucrose, xanthan gum

#### Questions or comments?

1-800-719-9260

### **Principal Display Panel - Daytime**

TWIN PACK

Compare to the active ingredients in Vicks® DayQuil® Severe

Severe

Non-Drowsy

Daytime

**COLD & FLU RELIEF** 

ACETAMINOPHEN – Pain reliever; Fever reducer

DEXTROMETHORPHAN HBr – Cough suppressant

**GUAIFENESIN** – Expectorant

PHENYLEPHRINE HCl – Nasal decongestant

MAXIMUM STRENGTH

Relieves:

Aches; Fever; Cough; Nasal congestion; Sore throat; Chest congestion

Alcohol free

Antihistamine free

Compare to the active ingredients in Vicks® NyQuil® Severe

Severe

Nighttime

COLD & FLU RELIEF

ACETAMINOPHEN – Pain reliever; Fever reducer

DEXTROMETHORPHAN HBr – Cough suppressant

DOXYLAMINE SUCCINATE – Antihistamine

PHENYLEPHRINE HCl - Nasal decongestant

MAXIMUM STRENGTH

Relieves:

Aches; Fever; Sore throat; Cough; Sneezing; Runny nose; Nasal/sinus congestion & Sinus pressure

Berry Flavor

Alcohol free

TWO 12 FL OZ (355 mL) UNITS TOTAL 24 FL OZ (1.5 PT) (710 mL)

www.StopMedicineAbuse.org

"These products are not manufactured or distributed by Procter & Gamble, distributor of Vicks® bayQuil® Severe and Vicks® by Quil® Severe.

Drug Facts (continued)

Inactive ingredients butylated hydroxyanisole, edetate disodium, FD&C yellow #5, flavor, glycerin, menthol, monobasic sodium phosphate polyethylene glycol, propylene glycol, purified water, saccharin sodium,

estions or comments?1-800-719-9260

**♥CVS**Health.

## TWIN PACK

Severe

Compare to the active ingredients in Vicks® DayQuil® Severe

**CVS** Health. Compare to the active ingredients in Vicks® NyQuil® Severe\*

Severe

NDC 59779-597-02

**Non-Drowsy** 

ACETAMINOPHEN - Pain reliever; Fever reducer **DEXTROMETHORPHAN HBr** - Cough suppressant **GUAIFENESIN** - Expectorant PHENYLEPHRINE HCI - Nasal decongestant

#### MAXIMUM STRENGTH

Relieves: Aches; Fever; Cough; Nasal congestion;



Alcohol free Antihistamine free

TWO 12 FL OZ (355 mL) UNITS

### **Nighttime COLD & FLU RELIEF**

ACETAMINOPHEN - Pain reliever; Fever reducer DEXTROMETHORPHAN HBr - Cough suppressant **DOXYLAMINE SUCCINATE - Antihistamine** PHENYLEPHRINE HCI - Nasal decongestant

#### MAXIMUM STRENGTH

#### Relieves:

Aches; Fever; Sore throat; Cough; Sneezing; Runny nose; Nasal/sinus congestion & Sinus pressure



**Berry Flavor** Alcohol free

TOTAL 24 FL OZ (1.5 PT) (710 mL)

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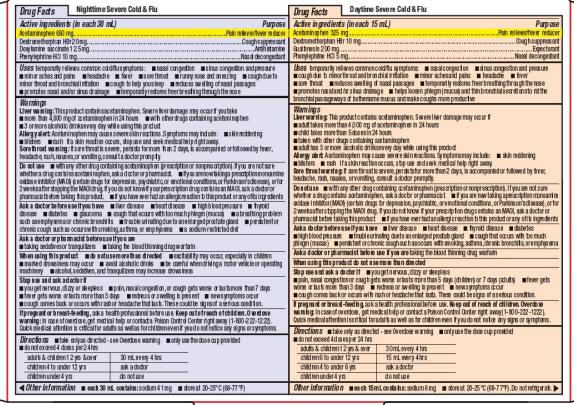
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nents?1-800-719-9260

Inactive ingredients antydrous citic acti, actate descrium, FD&C bue #1, FD&C red #40, flavor, glycefin, propleme glycol, purified water, seccharin sodium, sodium berzoate, sodium chloride, sodium dit ate, sorbibal solution, sucratose, xanthan gum

Drug Facts (continued)





#### SEVERE NIGHTTIME SEVERE DAYTIME

acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl kit

# Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:59779-597

l	Packaging				
	# Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
l	1 NDC:59779-597-02	1 in 1 CARTON; Type 0: Not a Combination Product	12/27/2013		

Quant	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

#### **SEVERE NIGHTTIME**

acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl solution

#### **Product Information**

Item Code (Source) NDC:59779-763

Route of Administration ORAL

#### **Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength 650 mg ACETAMINO PHEN (UNII: 36209 ITL9D) (ACETAMINO PHEN - UNII: 36209 ITL9D) ACETAMINOPHEN in 30 mL **DEXTROMETHO RPHAN HYDRO BRO MIDE** (UNII: 9 D2RTI9 KYH) **DEXTROMETHORPHAN** 20 mg (DEXTROMETHORPHAN - UNII:7355X3ROTS) **HYDROBROMIDE** in $30\ mL$ DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE -12.5 mg DOXYLAMINE SUCCINATE UNII:95QB77JKPL) in 30 mL PHENYLEPHRINE HYDRO CHLO RIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE -PHENYLEPHRINE 10 mg UNII:1WS297W6MV) HYDROCHLORIDE in 30 mL

Inactive Ingredients			
Ingredient Name	Strength		
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)			
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
FD&C RED NO. 40 (UNII: WZB9127XOA)			
GLYCERIN (UNII: PDC6A3C0OX)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			
SACCHARIN SODIUM (UNII: SB8ZUX40TY)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
SODIUM CHLORIDE (UNII: 451W47IQ8 X)			
SODIUM CITRATE (UNII: 1Q73Q2JULR)			
SORBITOL (UNII: 506T60A25R)			
SUCRALOSE (UNII: 96K6UQ3ZD4)			
XANTHAN GUM (UNII: TTV12P4NEE)			

Product Characteristics			
Color	RED (clear, dark)	Score	
Shape		Size	
Flavor	BERRY	Imprint Code	
Contains			

Ш	Pac	kaging			
	#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
	1 NI	DC:59779-763-40	355 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Markeniig iiiivi iiiauvii	Marl	keting	Information	1
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п				
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	OTC monograph final	part341	12/27/2013	

### Part 2 of 2

### **SEVERE DAYTIME**

acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl solution

### **Product Information**

Item Code (Source)	NDC:59779-603
Route of Administration	ORAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINO PHEN	325 mg in 15 mL		
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9 D2RTI9 KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 15 mL		
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 15 mL		
PHENYLEPHRINE HYDRO CHLO RIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 15 mL		

Inactive Ingredients		
Ingredient Name	Strength	
BUTYLATED HYDRO XYANISOLE (UNII: REK4960K2U)		
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)		
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)		
GLYCERIN (UNII: PDC6A3C0OX)		
MENTHOL (UNII: L7T10EIP3A)		
SODIUM PHO SPHATE, MO NO BASIC (UNII: 3980 JIH2SW)		
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SACCHARIN SO DIUM (UNII: SB8 ZUX40 TY)		
SUCROSE (UNII: C151H8 M554)		
XANTHAN GUM (UNII: TTV12P4NEE)		

Proc	luct	Chara	cteri	istics
1100	luct	Ciiai a		is tits

Color	ORANGE (clear)	Score	
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Shape		Size	
Flavor	FRUIT, MENTHOL	Imprint Code	
Contains			
Packaging			
# Item Code	Package Description	Marketing Start Date	<b>Marketing End Date</b>
1 NDC:59779-603-40	355 mL in 1 BOTTLE; Type 0: Not a Combination Produ	ct	
1 NDC:59779-603-40	355 mL in 1 BOTTLE; Type 0: Not a Combination Produ	ct	
1 NDC:59779-603-40  Marketing Inf		ct	
	ormation	Marketing Start Date	Marketing End Date
Marketing Inf	ormation		Marketing End Date
Marketing Inf	Ormation  Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Marketing Inf	Ormation  Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Marketing Inf	Ormation  Application Number or Monograph Citation  part341	Marketing Start Date	Marketing End Date

### Labeler - CVS Pharmacy (062312574)

part341

OTC monograph final

Revised: 1/2019 CVS Pharmacy

12/27/2013