# COOLING SEVERE DAYTIME COOLING SEVERE NIGHTTIME- acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride, guaifenes in 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. Cooling Severe Daytime Cooling Sever Nighttime Drug Facts

#### Active ingredients (in each 30 mL) - Night Time

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
- sore throat
- runny nose and sneezing
- cough to help you sleep
- cough due to minor throat and bronchial irritation
- reduces swelling of nasal passages
- fever
- 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
- be careful when driving a motor vehicle or operating machinery
- marked drowsiness may occur

- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic drinks

#### Stop use and ask a doctor if

- pain, nasal congestion, or cough gets worse or lasts more than 7 days
- you get nervous, dizzy or sleepless
- 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) - Day Time

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
- reduces swelling of nasal passages
- temporarily restores freer breathing through the nose
- sore throat
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- promotes nasal and/or sinus drainage

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- heart disease
- thyroid disease
- 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
- diabetes
- 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
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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

Compare to the active ingredients in  $Vicks^{\mathbb{R}}$  DayQuil $^{\mathbb{R}}$  Severe+VapoCOOL $^{\mathsf{TM}}$ 

Cooling

Severe Daytime

**COLD & FLU RELIEF** 

ACETAMINOPHEN - Pain reliever/Fever reducer

PHENYLEPHRINE HCl – Nasal decongestant

DEXTROMETHORPHAN HBr – Cough suppressant

**GUAIFENESIN** – Expectorant

Relieves:

Minor aches, pains & fever

Nasal congestion & sinus pressure

Cough

Chest congestion

**ALCOHOL 10%** 

12 FL OZ (355 mL)

Compare to the active ingredients in Vicks<sup>®</sup> NyQuil<sup>®</sup> Severe+VapoCOOL<sup>™</sup>

Cooling

Severe Nighttime

**COLD & FLU RELIEF** 

ACETAMINOPHEN – Pain reliever/Fever reducer

PHENYLEPHRINE HCl - Nasal decongestant

DOXYLAMINE SUCCINATE - Antihistamine

DEXTROMETHORPHAN HBr – Cough suppressant

Relieves:

Minor aches, pains & fever

Nasal congestion & sinus pressure

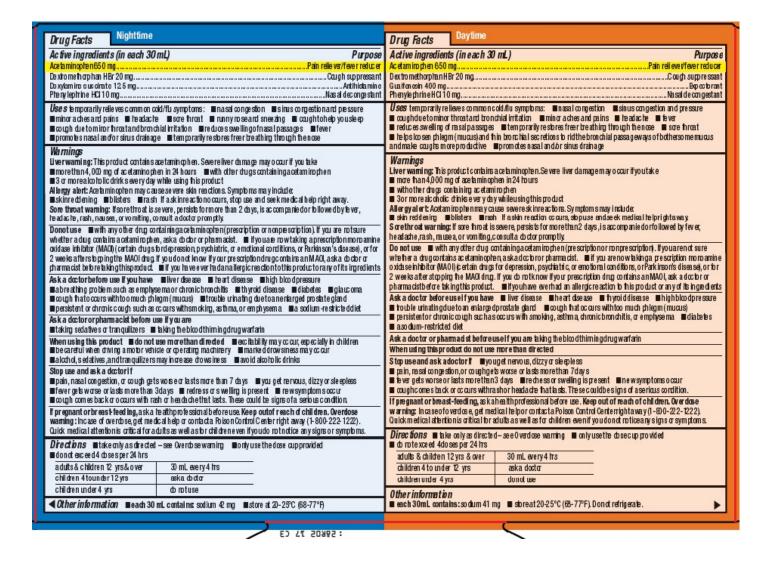
Sneezing, runny nose

Cough

**ALCOHOL 10%** 

12 FL OZ (355 mL)





#### COOLING SEVERE DAYTIME COOLING SEVERE NIGHTTIME

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

## Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:69842-737

	Packaging			
l	# Item Coc	le Package Description	Marketing Start Date	<b>Marketing End Date</b>
	1 NDC:69842-73	37-02 1 in 1 PACKAGE; Type 0: Not a Combina	tion Product 07/23/2019	

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

## **COOLING SEVERE NIGHTTIME**

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride solution

Product 1	Information
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Item Code (Source) NDC:69842-590

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)	ACETAMINOPHEN	650 mg in 30 mL	
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9 D2RTI9 KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 30 mL	
DO XYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DO XYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 30 mL	
PHENYLEPHRINE HYDRO CHLO RIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 30 mL	

Inactive Ingredients		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)		
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)		
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
GLYCERIN (UNII: PDC6A3C0OX)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SACCHARIN SO DIUM (UNII: SB8 ZUX40 TY)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)		
SORBITOL (UNII: 506T60A25R)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		

l	Packaging				
l	#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
l	1 N	DC:69842-590-40	355 mL in 1 BOTTLE; Type 0: Not a Combination Product		

	Marketing Information			
ı	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC monograph final part341 06/20/2019

## Part 2 of 2

## **COOLING SEVERE DAYTIME**

acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride solution

<b>Product Information</b>	
Item Code (Source)	NDC:69842-182
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)	ACETAMINOPHEN	650 mg in 30 mL	
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9 D2RTI9 KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 30 mL	
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 30 mL	
<b>PHENYLEPHRINE HYDRO CHLO RIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 30 mL	

Inactive Ingredients		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)		
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)		
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
GLYCERIN (UNII: PDC6A3C0OX)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
SO DIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)		
SORBITOL (UNII: 506T60A25R)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		

]	ackaging				
7	# Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
	NDC:69842-182-40	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 final	part341	06/20/2019				
Marketing Info	rmation					
Marketing Info	rmation Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
		Marke ting Start Date	Marketing End			

## Labeler - CVS Pharmacy (062312574)

Revised: 1/2020 CVS Pharmacy