

CVS DT AND NT COMBO PACK COLD AND FLU- acetaminophen, phenylephrine hydrochloride, and doxylamine succinate
CVS PHARMACY

CVS DT & NT Combo Pack Cold & Flu

CVS DayTime Cold & Flu Relief

Active ingredients (in each 15 mL)

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Phenylephrine HCl 5 mg

Purpose

Pain reliever/Fever reducer
Cough suppressant
Nasal decongestant

Uses

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

Warnings

Liver warning

This product contains acetaminophen.

Severe liver damage may occur if you take

- adult takes more than 4 doses (30 mL each) in 24 hours, which is the maximum daily amount for this product
- child takes more than 4 doses (15 mL each) in 24 hours, which is the maximum daily amount for this product
- 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.

Ask a doctor before use if you have

- Liver disease
- Heart disease
- High blood pressure
- Thyroid disease
- Diabetes
- 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
- 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 5 days (children) or 7 days (adult)
 - 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. In case of overdose, get medical help or contact a Poison Control Center right away. 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
- Only use the dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over	30 mL every 4 hours
Children 6 to under 12 yrs	15 mL every 4 hours
children 4 to under 12 yrs	ask a doctor
children under 4 yrs	do not use

Other information

each 15 mL contains:

- sodium 42 mg
- store at room temperature
- Do not refrigerate.

Inactive ingredients

Anhydrous citric acid, disodium edetate, FD&C Yellow No. 6, flavor, glycerin, menthol, potassium citrate, propyl gallate, purified water, sodium benzoate, sodium chloride, sorbitol, sucralose, xanthan gum

Questions?

1-866-467-2748

CVSNIGHTTIME COLD & FLU RELIEF

Active ingredients (in each 30 mL)

Acetaminophen 650 mg
Dextromethorphan HBr 30 mg
Doxylamine succinate 12.5 mg

Purpose

Pain reliever/Fever reducer
Cough suppressant
Antihistamine

Uses

temporarily relieves common cold/flu symptoms:

- cough due to minor throat & bronchial irritation
- minor aches & pains
- headache
- fever
- sore throat
- runny nose & 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
- taken 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.

Ask a doctor before use if you have

- Liver disease
- glaucoma
- Trouble urinating due to enlarged prostate gland
- Cough that occurs with too much phlegm (mucus)
- A breathing problem or chronic cough such as occurs with smoking, asthma, or emphysema

Ask a doctor or pharmacist before use if you are

- Taking sedatives or tranquilizers
- Taking the blood thinning drug warfarin

When using this product

- 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 & 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. In case of overdose, get medical help or contact a Poison Control Center right away. 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
- Only use the dose cup provided
- do not exceed 4 doses per 24 hrs

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

Other information

- each 30 mL contains: sodium 16 mg
- Store at room temperature

- Do not refrigerate

Inactive ingredients

Citric acid, D&C Yellow No. 10, disodium edetate, FD&C Green No.3, FD&C Yellow no 6, flavor, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose

Questions?

1-866-467-2748

Distributed By:**PRINCIPAL DISPLAY PANEL - Kit Carton**

Compare to the active ingredients in Vicks® DayQuil™ Cold & Flu*

NDC# 51316-618-12

DayTime**Cold & Flu****Relief**

Acetaminophen, Dextromethorphan HBr,

Phenylephrine HCl

- Headache, Fever, Sore Throat, Minor Aches & Pains
- Nasal Congestion & Sinus Pressure
- Cough

Compare to the active ingredients in Vicks® NyQuil™ Cold & Flu*

NightTime**Cold & Flu****Relief**

Acetaminophen, Dextromethorphan HBr

Dextromethorphan HBr

- headache, Fever, Sore Throat,
- Minor Aches & Pains
- Sneezing, Runny Nose
- Cough

Original Flavor

Tamper Evident: Do not use if imprinted shrink band is missing or broken.

TWO BOTTLES, 12 FL OZ (354 mL) each; TOTAL 24 FL OZ (708 mL)

***These product is not manufactured or distributed by Procter & Gamble, distributor of Vicks® Dayquil™ and Vicks® Nyquil™ Cold & flu.**



CVS DT AND NT COMBO PACK COLD AND FLU

acetaminophen, phenylephrine hydrochloride, and doxylamine succinate kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51316-618
---------------------	----------------	---------------------------	---------------

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51316-618-12	1 in 1 PACKAGE; Type 0: Not a Combination Product	03/20/2023	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	354 mL
Part 2	1 BOTTLE	354 mL

Part 1 of 2

CVS DAYTIME COLD AND FLU RELIEF

acetaminophen, dextromethorphan hydrobromide, and phenylephrine hydrochloride liquid

Product Information

Item Code (Source)	NDC:51316-620
---------------------------	---------------

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg in 15 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 15 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51316-620-12	354 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 Drug	M012	03/20/2023	

Part 2 of 2

CVS NIGHTTIME COLD AND FLU RELIEF

acetaminophen, doxylamine succinate, and dextromethorphan hydrobromide liquid

Product Information	
Item Code (Source)	NDC:51316-619
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg in 30 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 30 mL
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 30 mL

Inactive Ingredients	
Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51316-619-12	354 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 Drug	M010	03/20/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/20/2023	

Labeler - CVS PHARMACY (062312574)

Revised: 11/2023

CVS PHARMACY