DAYTIME NIGHTTIME COLD AND COUGH- acetaminophen, dextromethorphan hbr, phenylephrine hcl, doxylamine succinate
Walgreens

Drug Facts

Active ingredients (in each 15 mL) Daytime

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Phenylephrine HCl 5 mg

Active ingredients for (in each 30 mL) Nighttime

Acetaminophen 650 mg

Dextromethorphan HBr 30 mg

Doxylamine succinate 12.5 mg

Purposes for Day Time

Pain reliever/Fever reducer

Cough suppressant

Nasal decongestant

Purpose for Night Time

Pain reliever/fever reducer

Cough suppressant

Antihistamine

Uses

DAYTIME

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

NIGHTTIME

- temporarily relieves common cold/flu symptoms
  - minor aches and pain
  - headache
  - sore throat
  - runny nose and sneezing
  - fever
  - cough due to minor throat and bronchial irritations as may occur with a cold

Warnings

DAYTIME NIGHTTIME

Liver warning: This product contain acetaminophen. Severe liver damage may occur if:
- adult takes more than 4 doses (30 mL each) of acetaminophen in 24 hours, which is the maximum daily amount
- child takes more than 4 doses (15 mL each) 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

DAYTIME NIGHTTIME
- 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 contain an MAOI, ask a doctor or pharmacist before taking this product

Ask a doctor before use if have

DAYTIME
- liver disease
- high blood pressure
- heart disease
- thyroid disease
- diabetes
- a sodium-restricted diet

- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema
- cough that occurs with too much phlegm (mucus)

NIGHTTIME
- liver disease
- glaucoma
- cough that occurs with too much phlegm (mucus)
- trouble urinating due to enlarged prostate gland
- a breathing problem or chronic cough that lasts or occurs with smoking, asthma, chronic bronchitis, or emphysema

Ask a doctor or pharmacist before use if the child is

DAYTIME

if you are taking the blood thinning drug warfarin

NIGHTTIME
- taking sedatives or tranquilizers
- taking the blood thinning drug warfarin

When using this product

DAYTIME

do not exceed recommended dosage.

NIGHTTIME
- excitability may occur, especially in children
- mark 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

DAYTIME
- nervousness, dizziness or sleeplessness occur
- 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.persists for more than 7 days, comes back or occurs with a fever, rash or persistent headache.

These could be signs of a serious condition.

NIGHTTIME
- pain or cough gets worse or lasts more than 7 days
- redness or swelling is present
- new symptoms occur

- fever gets worse or lasts more than 3 days
- 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 the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Quick medical attention is critical for adult as well as for children even if you do not notice any signs or symptoms.

Directions

DAYTIME

- **do not take more than directed (see Overdose warning)**
- do not take more than 4 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other dosing device.
- keep dosing cup with product
- mL= milliliter

adult and children 12 years and over	30 mL every 4 hours
children 6 to under 12 years	15 mL every 4 hours
children 4 to under 6 years	ask a doctor
children under 4 years	do not use

NIGHTTIME

- **do not take more than directed (see Overdose warning)**
- do not take more than 4 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other dosing device.
- keep dosing cup with product
- mL= milliliter
- adult and children 12 years and over: 30 mL every 6 hours
- children under 12 years of age: do not use

Other information

DAYTIME

- **each 15 mL contains:** sodium 12 mg
- store between 20-25ºC (68º-77ºF). Do not refrigerate.

NIGHTTIME

- **each 30 mL contains:** potassium 5 mg

- **each 30 mL contains:** sodium 19 mL
- store between 20-25ºC (68-77ºF). Do not refrigerate

Inactive ingredients

DAYTIME

citric acid, FD&C yellow #6, flavor, glycerin, propylene glycol, purified water, saccharin sodium, sodium benzoate, sodium chloride, sodium citrate, sorbitol, sucralose, xanthan gum

NIGHTTIME

acesulfame potassium, alcohol, anhydrous citric acid, D&C yellow #10 FD&C green #3, FD&C yellow #6, flavor, high fructose corn syrup, polyethylene glycol, propylene glycol, purified water, saccharin sodium, trisodium citrate dihydrate

Questions or comments?

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal Display Panel

Compare to the active ingredients in Vick® Dayquil® Cold & Flu & Vick® Nyquil® Cold & Flu††

DAYTIME • NON-DROWSY

Cold & Cough

ACETAMINOPHEN 325 mg / PAIN RELIEVER / FEVER REDUCER

DEXTROMETHORPHAN HBr 10 mg / COUGH SUPPRESSANT

PHENYLEPHRINE HCl 5 mg / NASAL DECONGESTANT

Multi-Symptom
- Relieves aches, fever,& sore throat, cough & nasal congestion
- For ages 6 years & over
- Alcohol free
- Antihistamine free

FL OZ (mL)

NIGHTTIME

Cold & Flu

ACETAMINOPHEN 650 mg / PAIN RELIEVER / FEVER REDUCER

DEXTROMETHORPHAN HBr 30 mg / COUGH SUPPRESSANT

DOXYLAMINE SUCCINATE 12.5 mg / ANTIHISTAMINE

Multi-Symptom
- Relieves headache, fever, sore throat, minor aches & pains, sneezing, runny nose &

cough
- For ages 12 years & over
- Nighttime relief
- ALCOHOL 10%

FL OZ (mL)

WHEN USING OTHER DAYTIME OR NIGHTTIME PRODUCTS, CAREFULLY READ EACH LABEL TO ENSURE CORRECT DOSING

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL AROUND DOSAGE CUP OR UNDER CAP IS BROKEN OR MISSING

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

††These products are not manufactured or distributed by The Procter & Gamble Company. Vicks®, DayQuil® and Nyquil® are registered trademarks of the Procter & Gamble Company.

DISTRIBUTED BY: WALGREEN CO.

200 WILMOT RD., DEERFIELD, IL 60015 walgreens.com

Product Label

WALGREENS Daytime Nighttime Cold and Flu

DAYTIME NIGHTTIME COLD AND COUGH

acetaminophen, dextromethorphan hbr, phenylephrine hcl, doxylamine succinate kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-4670

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-4670-24	1 in 1 KIT; Type 0: Not a Combination Product	07/30/2020	02/28/2027

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

COLD AND FLU RELIEF

acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid

Product Information

Item Code (Source)	NDC:0363-4661
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)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
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		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 Drug	M012	07/30/2020	02/28/2027

Part 2 of 2

COLD AND FLU NIGHTTIME

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate liquid

Product Information	
Item Code (Source)	NDC:0363-3430
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
ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
ALCOHOL (UNII: 3K9958V90M)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	

SACCHARIN SODIUM (UNII: SB8ZUX40TY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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 Drug	M012	07/30/2020	02/28/2027

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/30/2020	02/28/2027

Labeler - Walgreens (008965063)

Revised: 9/2025 Walgreens