

COLD AND FLU RELIEF DAYTIME- acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride capsule, liquid filled H E B

Drug Facts

Active ingredients (in each softgel)

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 and flu symptoms:
 - minor aches and pains
 - headache
 - sore throat
 - nasal congestion
 - fever
 - cough due to minor throat and bronchial irritation

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.

Ask a doctor before use if you have

- liver disease
- diabetes
- heart disease
- thyroid disease
- high blood pressure
- 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)

Ask a doctor or pharmacist before use if you are

taking the blood thinning drug warfarin.

When using this product,

do not exceed recommended dosage.

Stop use and ask a doctor if

- pain, cough, or nasal congestion gets worse or lasts more than 7 days
- new symptoms occur
- nervousness, dizziness or sleeplessness occur
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- 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 can 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 adults as well as for children even if you do not notice any signs or symptoms.

Directions

- **do not take more than directed (see Overdose warning)**
- do not take more than 4 doses in 24 hours
- adults and children 12 years and over: take 2 softgels with water every 4 hours
- swallow whole; do not crush, chew, or dissolve
- children under 12 years: do not use
- **when using other Daytime or Nighttime products, carefully read each label to ensure correct dosing**

Other information

- store between 15-30°C (59-86°F)
- avoid excessive heat

Inactive ingredients

butylated hydroxyanisole, butylated hydroxytoluene, FD&C yellow #6, gelatin, glycerin, mannitol, polyethylene glycol, povidone, propylene glycol, purified water, sorbitan, sorbitol, white ink

*may contain this ingredient

Questions or comments?

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

Principal Display Panel

Compare to Vicks® DayQuil® Cold & Flu active ingredients**

Daytime

Acetaminophen / Pain Reliever / Fever Reducer

Dextromethorphan HBr / Cough Suppressant

Phenylephrine HCl / Nasal Decongestant

Cold & Flu

Non-Drowsy

Multi-Symptom

Relief of:

- Pain
- Fever
- Cough
- Allergy Symptoms

SOFTGELS†

(†LIQUID-FILLED CAPSULES)

**This product is not manufactured or distributed by The Procter & Gamble Company, Vicks® DayQuil®, are registered trademarks of The Procter & Gamble Company.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

MADE WITH PRIDE AND CARE FOR H-E-B®, SAN ANTONIO, TX 78204

Product Label



actual size

16 SOFTGELS[†]
(LIQUID-FILLED CAPSULES)

Relief of:
Multi-Symptom
Non-Drowsy
• Pain • Fever • Cough
• Nasal Congestion

Cold & Flu

Acetaminophen / Pain Reliever/Fever Reducer
Dextromethorphan HBr / Cough Suppressant
Phenylephrine HCl / Nasal Decongestant

Daytime



NDC 37808-848-16

Compare to Vicks® Dayquil® Cold & Flu active ingredients**

Drug Facts		Drug Facts (continued)	
Active ingredients (in each softgel)	Purposes	Uses	
Acetaminophen 325 mg.....	Pain reliever/fever reducer	■ temporarily relieves common cold and flu symptoms:	
Dextromethorphan HBr 10 mg.....	Cough suppressant	■ minor aches and pains	■ headache
Phenylephrine HCl 5 mg.....	Nasal decongestant	■ nasal congestion	■ fever
		■ cough due to minor throat and bronchial irritation	

Drug Facts (continued)

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.

Ask a doctor before use if you have ■ liver disease ■ diabetes
 ■ heart disease ■ thyroid disease ■ high blood pressure
 ■ 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)

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin.

When using this product, do not exceed recommended dosage.

Drug Facts (continued)

Stop use and ask a doctor if
 ■ pain, cough, or nasal congestion gets worse or lasts more than 7 days
 ■ nervousness, dizziness, or sleeplessness occur
 ■ 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: Taking more than the recommended dose can 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 adults as well as for children even if you do not notice any signs or symptoms.

Directions ■ do not take more than directed (see Overdose warning)
 ■ do not take more than 4 doses in 24 hours
 ■ adults and children 12 years and over: take 2 softgels with water every 4 hours
 ■ swallow whole; do not crush, chew, or dissolve
 ■ children under 12 years: do not use
 ■ when using other Daytime or Nighttime products, carefully read each label to ensure correct dosing

Other information
 ■ store between 15-30°C (59-86°F) ■ avoid excessive heat

Inactive ingredients butylated hydroxyanisole, butylated hydroxytoluene, FD&C yellow #6, gelatin, glycerin, mannitol, polyethylene glycol, povidone, propylene glycol, purified water, sorbitan, sorbitol, white ink

Questions or comments?
 Call toll free: 1-877-753-3935 Monday-Friday 9AM-5PM EST

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.
 KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Lot No.:
Exp. Date:

MADE WITH PRIDE AND CARE FOR H-E-B®
SAN ANTONIO, TX 78204

This product is not manufactured or distributed by The Procter
& Gamble Company. Vicks® and DayQuil® are registered
trademarks of The Procter & Gamble Company.

15989-2007



PLD-E5688 FC006076



H-E-B Daytime Cold and Flu

COLD AND FLU RELIEF DAYTIME

acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-848
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITAN (UNII: 6O92ICV9RU)	
SORBITOL (UNII: 506T60A25R)	
MANNITOL (UNII: 3OWL53L36A)	

Product Characteristics

Color	orange	Score	no score
Shape	CAPSULE	Size	20mm
Flavor		Imprint Code	P19

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-848-16	16 in 1 CARTON	06/30/2020	
1		1 in 1 BLISTER PACK; 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	06/30/2020	

Labeler - H E B (007924756)

Revised: 4/2024

H E B