COLD AND FLU DAYTIME- acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride capsule, liquid filled Safeway, Inc.

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 & 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 is 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
- nervousness, dizziness or sleeplessness occur
- fever gets worse or last 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 serious liver damage. In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical for adults and 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 insure 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 **1-888-723-3929** Monday- Friday 7AM- 6PM PST

Principal Display Panel

Compare to Vicks® DayQuil® Cold & Flu LiquiCaps® active ingredients†

Non-Drowsy Daytime

Cold & Flu Relief

ACETAMINOPHEN 325mg - Pain Reliever/Fever Reducer

DEXTROMETHORPHAN HBr 10 mg - Cough Suppressant

PHENYLEPHRINE HCL 5mg - Nasal Decongestant

- Non-drowsy
- Alcohol-free
- Antihistamine-free

SOFTGELS**

(**Liquid-filled capsules)

†This product is not manufactured or distributed by The Procter & Gamble Company, Vicks®, DayQuil®, and LiquiCaps® 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.

DISTRIBUTED BY: BETTER LIVING BRANDS LLC P.O. BOX 99, PLEASANTON, CA 94566-0009 www.betterlivingbrandsLLC.com

Product Label

When using this product, do not exceed recommended dosage.

типпппд огид warrarm.

Ask a doctor or pharmacist before use if you are taking the blood

- condu tust occurs with too much phiegm (mucus)
- becaistent or chronic cough such as occurs with smoking, asthma, or ■ trouble unnating due to an enfarged prostate gland
- high blood pressure ■ heart disease ■ thyroid disease

Ask a doctor before use if you have ■ liver disease ■ diabetes

bysrmacist before taking this product. do not know if your prescription drug contains an MAOI, ask a doctor or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you (certain drugs for depression, psychiatric, or emotional conditions, or

 if you are now taking a prescription monoamine oxidase inhibitor (MAOI) acetaminophen, ask a doctor or pharmacist.

uoublescubpou). It you are not sure whether a drug contains with any other drug containing acetaminophen (prescription or

vomiting, consult a doctor promptly.

days, is accompanied or followed by fever, headache, rash, nausea, or Sore throat warning: If sore throat is severe, persists for more than 2 If a skin reaction occurs, stop use and seek medical help right away. wsy include: ■ skin reddening ■ blisters ■ rash

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms

- 3 or more alcoholic drinks every day while using this product
 - with other drugs containing acetaminophen
 - more than 4,000 mg of acelaminophen in 24 hours may occur if you take:

Liver warning: This product contains acetaminophen. Severe liver damage Warnings

Drug Facts (continued)

C9II 1-811-123-3932 Woudsh-Friday 9AM-5PM EST Questions or comments?

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

> ■ store between 15-30°C (59-86°F)
> ■ avoid excessive heat попетиоти тэнтО

label to ensure correct dosing

- when using other Daytime or Nighttime products, carefully read each ■ children under 12 years: do not use

 - swallow whole; do not crush, chew, or dissolve
- adults and children 12 years and over: take 2 softgels with water every 4 hours ■ do not take more than 4 doses in 24 hours

DIRECTIONS

do not take more than directed (see Overdose warning)

adults as well as for children even if you do not notice any signs or symptoms. Center (1-800-222-1222) right away. Quick medical attention is critical for damage. In case of overdose, get medical help or contact a Poison Control Overdose warning: Taking more than the recommended dose can cause liver

Keep out of reach of children. If pregnant or breast-feeding, ask a health professional before use.

sidus ot a serious condition.

- cough comes back or occurs with rash or headache that lasts. These could be
 - Lequese or swelling is present
 Inew symptoms occur
 - fever gets worse or lasts more than 3 days
 - nervousness, dizziness, or sleeplessness occur ■ pain, cough, or nasal congestion gets worse or lasts more than 7 days
 - Stop use and ask a doctor if

cough due to minor throat and bronchial irritation

■ 16ver

■ headache ■ sore throat

Drug Facts (continued)

Nasal decongestant Phenylephrine HCl 5 mg. Congh suppressant Dextromethorphan HBr 10 mg. ALLI TEHEVET/TEVET TEDUCET Active ingredients (in each softgel) Purposes

NSGS # femborarily relieves common cold and flu symptoms:

■ winor aches and pains

(confined) **szzer** (confined)

■ usssi congestion

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

NDC 21130-650-24

Drug Facts



Non-Drowsy Daytime Cold & Flu Relief

ACETAMINOPHEN 325 mg - Pain Reliever/Fever Reducer

DEXTROMETHORPHAN HBr 10 mg - Cough Suppressant

PHENYLEPHRINE HCI 5 mg - Nasal Decongestant

Actual Size

24 SOFTGELS**

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

 Non-drowsy Alcohol-free DISTRIBUTED BY: BETTER LIVING BRANDS LLC P.O. BOX 99, PLEASANTON, CA 94566-0009 1-888-723-3929 ¹This product is not manufactured or distributed by The Procter & Gamble Company. Vicks®, DayQuit®, and LiquiCaps® are registered trademarks of The Procter & Gamble Company.

www.betterlivingbrandsLLC.com

PLD-H568B FC008030

OUR PROMISE
OUALITY & SATISFACTION
100% GUARANTEED
OR YOUR MONEY BACK.





Scan here for more information



SIGNATURE CARE Non Drowsy Daytime Cold & Flu Relief

COLD AND FLU DAYTIME

acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride capsule, liquid filled

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:21130-650

Route of Administration ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1W5297W6MV)	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: 6092ICV9RU)	
SORBITOL (UNII: 506T60A25R)	
MANNITOL (UNII: 30WL53L36A)	

Product Characteristics			
Color	orange	Score	no score
Shape	CAPSULE	Size	20mm

Flavor	Imprint Code	P19
Contains		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:21130-650- 24	24 in 1 CARTON	06/30/2019		
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	06/30/2019	

Labeler - Safeway, Inc. (009137209)

Revised: 5/2024 Safeway, Inc.