DAYTIME NIGHTTIME COLD FLU RELIEF- acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl Safeway, Inc.

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.

DRUG FACTS

Active ingredients for Daytime (in each softgel) Acetaminophen 325 mg

Dextromethorphan HBr 10 mg
Phenylephrine HCl 5 mg

Active ingredients for Nighttime (in each softgel)

Acetaminophen 325 mg

Dextromethorphan HBr 15 mg
Doxylamine succinate 6.25 mg

Purpose for Daytime

Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

Purpose for Nighttime

Pain reliever/fever reducer

Cough suppressant

Antihistamine

Uses

DAYTIME

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

- fever
- sore throat

NIGHTTIME

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

Warnings

DAYTIME

Liver warning: These products contain 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 these products

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.

NIGHTTIME

Liver warning: This product contain acetaminophen. Severe liver damage may occur if you take:

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- 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 following by fever, headache, rash, nausea, vomiting, consult a doctor promptly.

Do not use

DAYTIME

- 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.

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Ask a doctor before use if you have

DAYTIME

- liver disease
- heart disease
- diabetetes
- thyroid disease
- high blood pressure
- cough that occurs with too much phlegm (mucus)
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema

NIGHTTIME

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

Ask a doctor or pharmacist before use if you are

DAYTIME

taking the blood thinning drug warfarin

NIGHTTIME

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

When using this product,

DAYTIME

do not exceed recommended dosage

NIGHTTIME

- do not exceed recommended dosage
- excitability may occur, especially in children
- marked drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

DAYTIME

- nervousness, dizziness or sleeplessness occur
- pain, cough, and nasal congestion 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.

NIGHTTIME

- 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 headache that lasts.

These could be a 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

DAYTIME

- do not take more than directed (see Overdose warning)
- do not take more than 4 doses in 24 hours
- swallow whole; do not crush, chew, or dissolve
- adults and children 12 years and over; take 2 softgels with water every 4 hours.

• children under 12 years: do not use

NIGHTTIME

- do not take more than directed (see Overdose warning)
- do not take more than 4 doses in 24 hours
- swallow whole; do not crush, chew, or dissolve
- adults and children 12 years and over: take 2 softgels with water every 6 hours
- children under 12 years: do not use

Other information

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

Inactive ingredients

DAYTIME

FD&C yellow #6, FD&C Red #40, gelatin, glycerin, lecithin, light mineral oil, mannitol, polyethylene glycol, povidone, propylene glycol, purified water, sorbitan, sorbitol, white ink

NIGHTTIME

D&C yellow #10, FD&C blue #1, gelatin, glycerin, lecithin, light mineral oil, mannitol, polyethylene glycol, povidone, propylene glycol, purified water, sorbitan, sorbitol, white ink

Questions or comments?

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

Principal Display Panel

Compare to Vicks® Dayquil® Cold & Flu LiquiCaps® active ingredient†

Non-Drowsy Daytime

Cold & Flu Relief

ACETAMINOPHEN 325 mg - Pain Reliever/Fever Reducer

DEXTROMETHORPHAN HBr 10 mg - Cough Suppressant

PHENYLEPHRINE HCL 5 mg - Nasal Decongestant

Relieves

- Aches, Fever, Sore Throat
- Cough
- Nasal Congestion

Alcohol-Free

Antihistamine-Free

SOFTGELS**

(**Liquid-Filled Capsules)

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

NIGHTTIME

Cold & Flu Relief

ACETAMINOPHEN 325 mg - Pain Reliever/Fever Reducer

DEXTROMETHORPHAN HBr 15 mg - Cough Suppressant

DOXYLAMINE SUCCINATE 6.25 mg - Antihistamine

Relieves

- Aches, Fever, Sore Throat
- Cough
- Runny Nose, Sneezing
- Alcohol-Free

SOFTGELS

When using Daytime and Nighttime products, carefully read the labeling to ensure correct dosing

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

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOW 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

a children under 12 years; do not use

- m adults and children 12 years and over: take 2 softgels with water every
 - a swallow whole; do not crush, chew, or dissolve
 - m do not take more than 4 doses in 24 hours m do not take more than directed (see Overdose warning)

Directions

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modical attention is critical for adults as well as for children even if you do not help or contact a Poison Control Center (1-800-222-1222) right away. Quick recommended does can cause liver damage. In case of overdose, get medical keep out of reach of children. Overdose warning: Taking more than the If pregnant or breast-feeding, ask a health professional before use.

sibus of a senous condition.

- cough comes back or occurs with rash or headache that lasts. These could be
 - Leguese ot swelling is present
 new symptoms occur
 - fever gets worse or lasts more than 3 days
 - bain or cough gets worse or lasts more than 7 days
 - Stop use and ask a doctor if
 - pe cereful when driving a motor vehicle or operating machinery
 - sicohol, sedalives, and tranquilizers may increase drowsiness misured drowsiness may occur.
 svoid stcoholic drinks
 - When using this product to not exceed recommended doesge excitability may occur, especially in children

missing the blood thinning drug warfarin mataing sedatives or transport

Ask a doctor or pharmacist before use if you are

- m trouble uninsting due to an enlarged prostate gland asthma, chronic bronchitts, or emphysema
- a breathing problem or chronic cough that lasts such as occurs with smoking,
 - condly first occurs with too much phiegm (mucus)

Ask a doctor before use if you have as liver disease as glaucoma

Drug Facts (continued)

Nighttime Cold & Flu Softgel

children under 12 years: do not use

 adults and children 12 years and over: take 2 softgels with water every m swallow whole; do not crush, chew, or dissolve auton 45 ni assob 4 nent enom exist fon ob m

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When using this product, do not exceed recommended dosage.

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

sak a doctor or pharmacist.

persistent or chronic cough such as occurs with smoking, asthma,

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Drug Facts (continued) Daytime Cold & Flu Softgel

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with any other drug containing acetaminophen (prescription or

3 or more alcoholic drinks every day while using this product

occur if you take: minore than 4,000 mg of acetaminophen in 24 hours

Parkinson's disease), or for 2 weeks after stopping the MADI drug. If you do

nonprescription). If you are not sure whether a drug contains acetaminophen,

accompanied or followed by fever, headache, resh, nausea, or vomiting, consult

Sore throat warning: If sore throat is severe, persists for more than 2 days, is If a skin reaction occurs, stop use and seek medical help right away.

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

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If you are now taking a prescription monoamine exidase won eas uoy 11 ask a doctor or pharmacist. nonprescription). If you are not sure whether a drug contains acetaminophen,

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Allengy alent. Acetaminophen may cause severe skin reactions. Symptoms may 3 or more alcoholic drinks every day while using this product

■ with other drugs containing acetaminophen

Active ingredients (in each softgel)

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Bulzaeus & saon ymnn ■ ■ headache ■ minor aches and pains ■ fever cough due to minor throat and bronchial imfation some throat nzes . I temporarily relieves common cold and flus symptoms:

animatairthnA Doxylamine succinate 6.25 mg. Cough suppressant Dextromethorphan HBr 15 mg.

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Phenylephine HCl 5 mg. рехфолетокрая НВг 10 тр. Pain reliever/lever reducer

uciode: m skin reddening m bisters m rash

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Active ingredients (in each softgel)

TREES . I semborarily relieves common cold and flu symptoms.

Nighttime Cold & Flu Softgel

Drug Facts

Daytime Cold & Flu Softgel

Compare to Vicks NyQuil Cold & Flu LiquiCaps' active ingredients

NDC 21130-901-48

Drug Facts

Signature care.

Compare to Vicks" DayQuil" Cold & Flu LiquiCaps'

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Nighttime Cold & Flu Relief

Non-Drowsy **Daytime** Cold & Flu Relief



SIGNATURE CARE Daytime Nighttime Cold & Flu Relief

DAYTIME NIGHTTIME COLD FLU RELIEF

acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:21130-901

l	P	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1	NDC:21130-901- 48	1 in 1 KIT; Type 0: Not a Combination Product	12/27/2019			

Quantity	of Parts	
Part #	Package Quantity	Total Product Quantity

Part 1	16 BLISTER PACK	16
Part 2	32 BLISTER PACK	32

Part 1 of 2

NIGHTTIME COLD FLU RELIEF

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate capsule

Product Information

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	15 mg		
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg		

Inactive Ingredients				
Ingredient Name	Strength			
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
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: 3OWL53L36A)				
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)				
LIGHT MINERAL OIL (UNII: N6K5787QVP)				

Product Characteristics			
Color green Score		Score	no score
Shape	CAPSULE	Size	21mm
Flavor		Imprint Code	PC10
Contains			

Packaging

	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1		16 in 1 CARTON		
	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 final	part341	12/27/2019		

Part 2 of 2

DAYTIME COLD FLU RELIEF

acetaminophen, dextromethorphan hbr, phenylephrine hcl capsule

Product Information

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: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg	

Inactive Ingredients			
Ingredient Name	Strength		
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)			
FD&C RED NO. 40 (UNII: WZB9127XOA)			
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)			
LIGHT MINERAL OIL (UNII: N6K5787QVP)			

Product Characteristics			
Color orange Score no sc			no score
Shape	CAPSULE	Size	20mm
Flavor		Imprint Code	PC9
Contains			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1		32 in 1 CARTON				
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 final	part341	12/27/2019		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part341	12/27/2019		

Labeler - Safeway, Inc. (009137209)

Revised: 2/2022 Safeway, Inc.