DAYTIME NON DROWS COLD AND FLU AND NIGHTTIME COLD ANDacetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride Spirit Pharmaceuticals LLC

Daytime Non Drowsy Cold and Flu and Nighttime Cold and Flu

Active ingredients (in each softgel)

COLD & FLU NON-DROWSY DAY RELIEF

Acetaminophen 325 mg Dextromethorphan hydrobromide 10 mg Phenylephrine hydrochloride 5 mg

COLD & FLU NIGHT RELIEF

Acetaminophen 325 mg Dextromethorphan hydrobromide 10 mg

Doxylamine succinate 6.25 mg

Purposes

COLD & FLU NON DROWSY DAY RELIEF

Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

COLD & FLU NIGHT RELIEF

Pain reliever/fever reducer

Cough suppressant

Antihistamine

Uses

temporarily relieves common cold/flu symptoms:

- fever
- headache
- minor aches and pain
- cough due to minor throat and bronchial iffitation
- sore throat
- nasal congestion (Daytime only)
- runny nose and sneezing (Nighttime only)

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 ● 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: \bullet skin reddening \bullet blisters \bullet 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.● if you have ever had an allergic reaction to this product or any of its ingredients● to make a child sleepy (Nighttime only)

Ask a doctor before use if you have

- ullet cough that occurs with too much phlegm (mucus) ullet liver disease
- trouble urinating due to enlarged prostate gland
- diabetes (Daytime only) heart disease (Daytime only)
- thyroid disease (Daytime only) high blood pressure (Daytime only)
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema (Daytime only)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis, or emphysema (Nighttime only)
- glaucoma (Nighttime only)

Ask a doctor or pharmacist before use if you are

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers (Nighttime only)

When using this product

- do not take more than directed
- marked drowsiness may occur (Nighttime only)
- avoid alcoholic drinks (Nighttime only)
- excitability may occur, especially in children (Nighttime only)
- be careful when driving a motor vehicle or operating machinery (Nighttime only)
- alcohol, sedatives, and tranquilizers may increase drowsiness (Nighttime only)

Stop use and ask a doctor if

- you get nervous, dizzy or sleepless (Daytime only)
- pain, nasal congestion, or cough gets worse or lasts more than 7 days (Daytime only)
- pain or cough gets worse or lasts more than 7 days (Nighttime only)
- 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 directed can cause serious health problems. In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical for adults & for children even if you do not notice any signs or symptoms.

Directions

• when using other DAYTIME and NIGHTTIME products, carefully read each label to ensure correct dosing

Directions (Daytime only)

- take only as directed see Overdose warning
- do not exceed 4 doses per 24 hours

adults & children 12 years & over	take 2 softgels with water every 4 hours
children 4 to under 12 years	ask a doctor
children under 4 years of age	do not use

• when using other DAYTIME and NIGHTTIME products, carefully read each label to ensure correct dosing

Directions (Nighttime only)

- take only as directed see Overdose warning
- do not exceed 4 doses per 24 hours

adults & children 12 years & over	take 2 softgels with water every 6 hours
children 4 to under 12 years	ask a doctor
children under 4 years of age	do not use

Other information

• store at room temperature.

Inactive ingredients

DAYTIME

FD&C Red# 40, FD&C Yellow# 6, gelatin, glycerin, polyethylene glycol, lecithin, myglyol, povidone, propylene glycol, purified water, sorbitol sorbitan, titanium dioxide

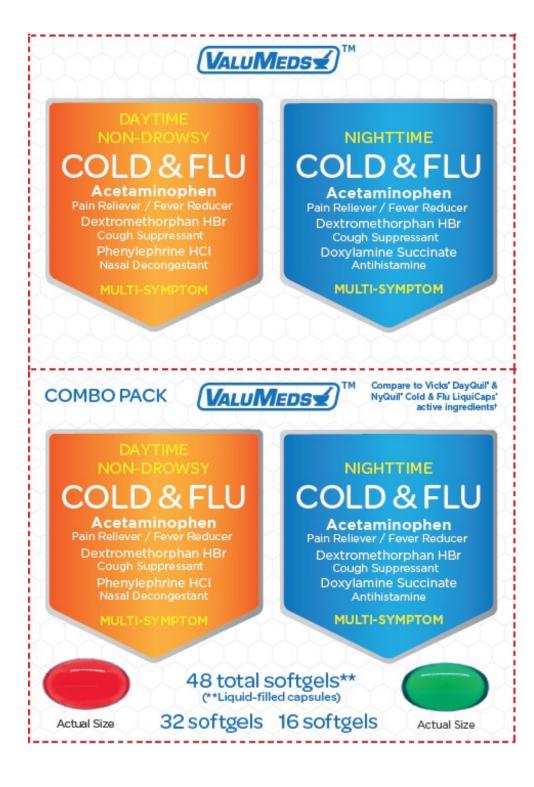
NIGHTTIME

D&C Yellow# 10, FD&C Blue# 1, gelatin, glycerin, myglyol, lecithin, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol sorbitan, polysorb, sorbitol sorbitan, titanium dioxide

Questions or comments?

1-888-333-9792

Principal Display Panel



Drug Facts	Drug Facts (continued)
Active ingredients for Daytime (in each softgel) Pain reliever.Never reducer Acetaminophen 325 mg Pain reliever.Never reducer Dextromethorphan HBr 10 mg Cough suppressant Phenylephrine HCI 5 mg Nacal decongestant Active ingredients for Nighttime Purpose (in each softgel) Active ingredients for Nighttime Purpose Active ingredients for Nighttime Purpose (in each softgel) Acetaminophen 325 mg Pain reliever.Never reducer Dextromethorphan HBr 15 mg Cough suppressant Doxylamine succinate 0.25 mg Dextromethorphan HBr 15 mg Cough suppressant Doxylamine succinate 0.25 mg Hempocarily relieves common cold' flu symptoms: Ever I headache II sore threat II minor aches and pains Bough due to minor threat and bronchial irritation Inacal congestion (Dayftime only) Warnings I'uver warning This product contains acetaminophen. Severe liver damage may occur if you take: Iwrore than 4 doces in 24 hours, which is the maximum daily amount for this product Mallergy alark t. Acetaminophen may cause severe skin reactions. Symptoms may include: Bor more alcoholic dinks every day while using this product Mallergy instruct. Acetaminophen may cause severe skin reactions. Symptoms may include:	Do not use I with any other drug containing acetaminophen (prescription or nonpeccription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmasist. If you are now taking a prescription monoamine oxidase inhibitor (MAOI) (cortain drugs for depression, pcychiatric 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 pharmasits before taking this product. If you have ever had an allergic reaction to this product or any of its ingredients It o make a child sleepy (Nightime only) Ask a doctor before use if you have It omake a child sleepy (Nightime only) Ask a doctor before use if you have It ongh that course with noo much philogm (mucus) II liver disease It touche urinating due to enlarged proctate gland I diabeter (Defitime only) II hard disace (Defitime only) I thyroid disease (Defitime only) III hard disace (Defitime only) I thyroid disease (Defitime only) I thyroid disease (Defitime only) I thereathing problem or chronic eough that lasts or as occurs with smoking, asthma, chronic bronchitis, or emphysema (Mightime only) I glaucoma (Mightime only) Ask a doctor or pharmasist before use if you are I taking sedatives or tranquitizers (Mightime only) When using this product I do not take more than directed I marked drousiness may occur (Mightime only) I exitability may occur, especially in children (Mightime only) I exitability may occur, especially in children (Mightime only) I alcoholis chinks (Mightime only) I alcoholis drives (Mightime only) I alcoholis drives (Mightime only) I avoid alcoholis drives (Mightime only) I alcoholis drives (Mightime o
Drug Facts (continued)	Drug Facts (continued)
Stop use and ask a doctor if Stop use and ask a doctor if you get nervous, diczy or skepless (D'ayfime only) pain, nasal congestion, or cough gets worse or lasts more than 7 days (Dayfime only) pain or cough gets worse or lasts more than 7 days (Nightime only) lever gets worse or lasts more than 3 days lever gets worse of a serious condition. If pregnent or breast-faeding, ask a health professional before use. Keep out of reach of children. Overdiose warning Taking more than directed can cause serious health problems. In case of overdose, get medical help or contact a Poison Control Center right anay. Quick medical help or contact a Poison Control Center right anay. Quick medical help or contact a Poison Control Center right anay. Quick medical attention is critical for aduts & for children even if you do not notice any signs or symptoms.	Directions (Nighttime only) Take only as directed - see Overdose warning to onet exceed 4 doces per 24 hours adults and children 12 years and over: take 2 softgels with water every 6 hours thildren 4 to under 12 years: ask a doctor thildren under 4 years: do not use when using other DATTIME and NIGHTTIME products, carefully read each label to ensure correct docing Other information the store at room temperature Inactive ingredients (Daytime only) FD&C Red No. 40, FD&C Yellow No. 8, gelatin, glycerin, lecithin, myglyol polyethylene glycol, povione, progytene glycol, purified water, sorthol
Directions (Daytime only) E take only as directed - see Overdose warning I do not exceed 4 doces per 24 hours adults and children 12 years and over: take 2 softgels with water every 4 hours E children 4 to under 12 years: ask a doctor E children under 4 years: do not use when using other DAYTIME and MIGHTTIME products, carefully read	poryanyana gyca, portana, propyana gyca, parina water, commo sorbitan, fitanium dioxide Inactive ingredients (Nighttime only) D&C Vielow No. 10, FD&C Blue No. 1, gelatin, glycerin, lecithin, myglyol, polyethylene glycal, povidone, propylane glycal, purified water, sorbital sorbitan, fitanium dioxide Questions or comments? Call 1-888-333-9782

DAYTIME NON DROWS COLD AND FLU AND NIGHTTIME COLD AND

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride kit

Ρ	roduct Infor	mation			
Ρ	roduct Type	HUMAN OTC DRUG	Item Code	e (Source)	NDC:68210-5008
P	ackaging				
#	ltem Code	Package D	escription	Marketing Start Date	Marketing End Date
	Item Code NDC:68210- 5008-4	Package D 1 in 1 CARTON; Type 1: C Package		-	-

Quant	Quantity of Parts					
Part #	Package Quantity	Total Product Quantity				
Part 1	1 BLISTER PACK	8				
Part 2	1 BLISTER PACK	8				

Part 1 of 2

DAYTIME NON DROWSY COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride capsule, liquid filled

Product Information

ltem Code (Source)

Route of Administration

ORAL

NDC:68210-5006

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	5 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C RED NO. 40 (UNII: WZ B9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITOL (UNII: 506T60A25R)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	red	Score	no score
Shape	OVAL	Size	21mm
Flavor		Imprint Code	512;A09;AP01

Contains							
Packaging	l						
# Item Code		Pack	age Description	Mark	eting Start Date		ing End ate
1	4 in 1	CARTON					
1	8 in 1 Produc		K; Type 0: Not a Combination				
Marketi	n <mark>g I</mark> n	format	ion				
Marketi Catego		Applicat	tion Number or Monograph Citation	Ма	rketing Start Date		ting Enc ate
OTC Monograp	h Drug	M012		12/12	2/2022		
nydrochloric Product Ir			filled				
Item Code (NDC:68210-5007				
Route of Ad			ORAL				
Active Ing	redier	nt/Active	Moiety				
		Ingred	lient Name		Basis of St	rength	Streng
ACETAMINOP	HEN (U	NII: 36209ITL	9D) (ACETAMINOPHEN - UNII:362O	9ITL9D)	ACETAMINOPHEN		325 mg
DEXTROMETH DEXTROMETH			ROMIDE (UNII: 9D2RTI9KYH) 3ROTS)		DEXTROMETHORF HYDROBROMIDE	PHAN	10 mg
DOXYLAMINE JNII:95QB77JK		NATE (UNII: \	/9BI9B5YI2) (DOXYLAMINE -		DOXYLAMINE SUC	CINATE	6.25 mg
Inactive Ir	ngredi	ents					
			Ingredient Name			Str	ength
FD&C BLUE N			1BD)				
GELATIN (UNII							
GLYCERIN (UN							
LECITHIN, SO							
			(UNII: C9H2L21V7U)				
POLIEIHILE	NE GLY	COL 400 (UI	NII: B697894SGQ)				

POVIDONE (UNII: FZ989GH94E)

sn		OL (UNII: 6DC9Q167\	/3)			
50	RBITOL (UNII: 50)6T60A25R)				
so	RBITAN (UNII: 60	092ICV9RU)				
тіт	ANIUM DIOXIDI	E (UNII: 15FIX9V2JP)				
Pr	oduct Chara	acteristics				
Co	lor	green	Score		no score	
Sh	аре	OVAL	Size		21mm	
Fla	vor		Imprint Code		116;A07;A	202
Co	ntains					
Pa	ckaging					
#	ltem Code	Package	Description	Marketing Date		Marketing End Date
1	2 in	1 CARTON				
1		1 BLISTER PACK; Typ duct	pe 0: Not a Combination			
Μ	arketing	Information				
M	arketing Marketing Category		Number or Monograph Citation	Marketi Da		Marketing End Date
	Marketing	Application				Marketing End Date
ΟΤ	Marketing Category C Monograph Dru	Application		Da		-
ΟΤ	Marketing Category C Monograph Dru arketing	Application M012	Citation	Da 10/19/2022	te	Date
ΟΤ	Marketing Category C Monograph Dru	Application M012		Da	te ng Start	-

Labeler - Spirit Pharmaceuticals LLC (179621011)

Revised: 12/2023

Spirit Pharmaceuticals LLC