DAYTIME COLD AND FLU- acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride capsule, liquid filled Strive Pharmaceuticals 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.

RIGHT REMEDIES DayTime Cold & Flu

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/flu symptoms: \bullet nasal congestion \bullet cough due to minor throat & bronchial irritation \bullet sore throat \bullet headache \bullet minor aches & pains \bullet fever

Warnings

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

- more than 4 doses in 24 hrs, which is the maximum daily amount for this product
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks daily 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, lasts for more than 2 days, occurs with 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 • heart disease • high blood pressure • thyroid disease • diabetes • trouble urinating due to enlarged prostate gland • cough that occurs with too much phlegm (mucus) • persistent or chronic cough as occurs with smoking, asthma, or emphysema

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

When using this product, do not use more than directed.

Stop use and ask a doctor if • you get nervous, dizzy or sleepless

- pain, nasal congestion or cough get worse or last 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 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: In case of overdose, get medical help or contact a Poison Control Center right away.

Prompt medical attention is critical for adults as well as children even if you do not notice any signs or symptoms.

Directions

- take only as directed see overdose warning
- do not exceed 4 doses per 24 hrs

2 softgels with water
every 4 hrs
ask a doctor
do not use

Other information

store at room temperature between 20-25°C (68-77°F)

Inactive ingredients

edible white ink, FD&C red #40, FD&C yellow #6, gelatin, glycerin, polyethylene glycol 400, povidone, propylene glycol, purified water, sorbitol sorbitan solution

Questions or comments

1-888-577-8033 Monday - Friday 8am - 4pm EST

Compare to the active ingredients of **Vicks[®] DayQuil™** Cold & Flu Relief LiquiCaps[™]*

RIGHT REMEDIES

READ AND KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

*This product is not manufactured or distributed by Procter & Gamble Company, owner of the registered trademarks Vicks[®] and DayQuil[™].

Distributed by: Strive Pharmaceuticals Inc., East Brunswick, NJ 08816

Product of UAE

Packaged and Quality Assured in the USA

DO NOT USE IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

REV.00-072022

Packaging





DRUG FACTS TABLE

Drug Facts	
Active ingredients (in each softgel) Pa Acetaminophen 325 mgPain reliever/Fever Dextromethorphan HBr 10 mgCough suppi Phenylephrine HCl 5 mgNasal decon	reducer ressant gestant
Uses temporarily relieves common cold/flu symptoms: ■ nasal congestion ■ cough due to throat & bronchial irritation ■ sore throat ■ headache ■ minor aches & pains ■ f	
Warnings Liver warning: This product contains acetaminophen. Severe liver damage may occur if you more than 4 doses in 24 hrs, which is the maximum daily amount for this product with other drugs containing acetaminophen 3 or more alcoholic drinks daily while using this product Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include: skin readtion occurs, stop use and seek medical help right away. Sore throat warning: If sore throat is severe, lasts for more than 2 days, occurs with or is for by fever, headache, rash, nausea, or vomiting, consult a doctor prompty.	
Do not use ■ with any other drug containing acetaminophen (prescription or nonprescription) are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist ■ if you are taking a prescription monoamine oxidase inhibitor (MAOI), (certian drugs for depression, psychia emotional conditions, or Parkinson's disease), or for 2 weeks after stoping the MAOI drug. If you know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this provide the taking the taken of taken of the taken of the taken of the taken of the taken of taken	atric or do not
Ask a doctor before use if you have ■ liver disease ■ heart disease ■ high blood pressure ■ disease ■ diabetes ■ trouble urinating due to enlarged prostate gland ■ cough that occurs with the phlegm (mucus) ■ persistent or chronic cough as occurs with smoking, asthma, or emphysema	thyroid oo much
Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfar	in.
When using this product, do not use more than directed.	
Stop use and ask a doctor if ■ you get nervous, dizzy or sleepless ■ pain, nasal congestion or cough get worse or last 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 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: In case of overdose, get medical help or contact a Poison Control Center right a	way.

Prompt medical attention is critical for adults as well as children even if you do not notice any signs or symptoms.

Drug Facts (continued)

Directions

- take only as directed see overdose warning
- do not exceed 4 doses per 24 hrs

adults and children 12 yrs & over	2 softgels with water every 4 hrs
children 4 to under 12	ask a doctor
children under 4 vrs	do not use

Other information

store at room temperature between 20-25°C (68-77°F)

Inactive ingredients

edible white ink, FD&C red #40, FD&C yellow #6, gelatin, glycerin, polyethylene glycol 400, povidone, propylene glycol, purified water, sorbitol sorbitan solution

Questions or comments

1-888-577-8033 Monday - Friday 8am - 4pm EST

DAYTIME COLD AND FLU

acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride capsule, liquid filled

Product Infor	mation								
Product Type		HUMAN OTC	DRUG	Item C	ode (Source)	NDC:706	92-807	
Route of Admin	istration	ORAL							
Active Ingred	ient/Active	Moiety							
	Ingre	dient Nam	е			Basis of S	Strength	Strengt	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII: 36209ITL9D) ACETAMINOPHEN						EN	325 mg		
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH)DEXTROMETHORPHAN(DEXTROMETHORPHAN - UNII:7355X3ROTS)HYDROBROMIDE						10 mg			
PHENYLEPHRINE I UNII:1WS297W6MV)	HENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - PHENYLEPHRINE - NII: 1WS297W6MV) PHENYLEPHRINE - HYDROCHLORID					_	5 mg		
Inactive Ingre	edients								
		Ingredie	nt Name				St	Strength	
FD&C RED NO. 40	-	-							
FD&C YELLOW NO									
GELATIN, UNSPEC		G86QN327L)							
GLYCERIN (UNII: PI	· · ·								
POLYETHYLENE G									
POVIDONE, UNSP)						
PROPYLENE GLYC		JQ167V3)							
SORBITOL (UNII: 506T60A25R) SORBITAN (UNII: 6092ICV9RU)									
Product Chara	acteristics								
Color	re	ed	Score				no score		
Shape	0	VAL	Size				21mm	mm	
Flavor			Imprint Co	de			807	7	
Contains			-						
Packaging									
# Item Code	Pa	ackage Des	scription		Mar	keting Start Date		eting End Date	
1 NDC:70692- 807-48	4 in 1 CARTON	I			04/05/	2023			
1	12 in 1 BLISTE Product	R PACK; Type	0: Not a Com	bination					
Marketing	Informat	tion							
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Category Citation Date	Date
OTC monograph final part341 04/05/2023	

Labeler - Strive Pharmaceuticals Inc. (080028013)

Revised: 5/2023

Strive Pharmaceuticals Inc.