NON DROWSY DAYTIME AND NIGHTTIME SINUS CONGESTION AND COUGH RELIEF- acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride CVS PHARMACY, INC.

CVS Health Non Drowsy Daytime and Nighttime Sinus Congestion and Cough Relief

Non Drowsy DAYTIME Sinus Congestion and Cough Relief Active ingredients (in each softgel)

Acetaminophen 325 mg

Dextromethorphan hydrobromide 10 mg Phenylephrine hydrochloride 5 mg

NIGHTTIME Sinus Congestion and Cough Relief Active ingredients (in each softgel)

Acetaminophen 325 mg Dextromethorphan hydrobromide 10 mg Doxylamine succinate 6.25 mg Phenylephrine hydrochloride 5 mg

Purposes

Non Drowsy DAYTIME Sinus Congestion and Cough Relief

Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

Purposes

Nighttime Sinus Congestion and Cough Relief

Pain reliever/fever reducer

Cough suppressant

Antihistamine

Nasal decongestant

Uses

Non Drowsy DAYTIME Sinus Congestion and Cough Relief

• temporarily relieves these symptoms due to a cold or flu:

- minor aches and pains
- headache
- cough
- sore throat
- nasal and sinus congestion
- temporarily reduces fever

Uses

NIghttime Sinus Congestion and Cough Relief

- temporarily relieves these symptoms due to a cold or flu:
 - minor aches and pains
 - headache
 - nasal and sinus congestion
 - cough
 - sore throat
 - runny nose
 - sneezing
- temporarily reduces fever

Non Drowsy DAYTIME Sinus Congestion and Cough Relief *Warnings*

Liver warning

These products contain acetaminophen. Severe liver damage may occur if you take

- more than 4000 mg of Acetaminophen 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

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.

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.

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
- in children under 12 years of age

Ask a doctor before use if you have

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- cough with excessive phlegm (mucus)
- difficulty in urination due to enlargement of the prostate gland
- persistent or chronic cough such 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 exceed recommended dosage

Stop use and ask a doctor if

- pain, cough, or nasal congestion 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 or headache that lasts.

These could be signs of a serious condition.

nervousness, dizziness, or sleeplessness occurs

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Nighttime Sinus Congestion and Cough Relief

Warnings

Liver warning

These products contain acetaminophen. Severe liver damage may occur if you take

- more than 4000 mg of Acetaminophen 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

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.

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.

Do not use to sedate children.

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
- in children under 12 years of age

Ask a doctor before use if you have

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- glaucoma
- cough with excessive phlegm (mucus)
- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema

Ask a doctor or pharmacist before use if you are

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

When using this product

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

Stop use and ask a doctor if

• pain, cough, or nasal congestion 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 or headache that lasts.

These could be signs of a serious condition.

• nervousness, dizziness, or sleeplessness occurs

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Non Drowsy DAYTIME Sinus Congestion and Cough Relief Directions

- do not take more than the recommended dose
- adults and children 12 years and over: take 2 softgels with water every 4 hours. Do not exceed 10 softgels in 24 hours or as directed by a doctor.
- children under 12 years: do not use

Nighttime Sinus Congestion and Cough Relief Directions

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- adults and children 12 years and over: take 2 softgels with water every 4 hours. Do not exceed 10 softgels in 24 hours or as directed by a doctor.
- children under 12 years: do not use

Non Drowsy DAYTIME Sinus Congestion and Cough Relief Other information

• store at room temperature. Avoid excessive heat.

Nighttime Sinus Congestion and Cough Relief Other information

• store at room temperature. Avoid excessive heat.

Non Drowsy DAYTIME Sinus Congestion and Cough Relief Inactive ingredients

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

Nighttime Sinus Congestion and Cough Relief Inactive ingredients

D&C Yellow No. 10, FD&C Blue No. 1, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol sorbitan, sodium hydroxide†, titanium dioxide †may contain this ingredient

Non Drowsy DAYTIME Sinus Congestion and Cough Relief

Questions or comments?1-888-333-9792

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PRINCIPAL DISPLAY PANEL



Compare to the active ingredients in Alka-Seltzer PLUS® Maximum Strength Severe Sinus Congestion & Cough Day & Night Liquid Gels*

Sinus Congestion & Cough

Daytime Non-Drowsy

Sinus Congestion & Cough Relief

ACETAMINOPHEN 325 mg Pain reliever/Fever reducer DEXTROMETHORPHAN HBr 10 mg Cough suppressant PHENYLEPHRINE HCI 5 mg Nasal decongestant

Relieves:

Actual Size

12 SOFTGELS

Nasal & sinus congestion; Sinus pressure; Headache & body ache; Cough

Nighttime Sinus Congestion & Cough Relief

ACETAMINOPHEN 325 mg Pain reliever/Fever reducer DEXTROMETHORPHAN HBr 10 mg Cough suppressant DOXYLAMINE SUCCINATE 6.25 mg Antihistamine PHENYLEPHRINE HCI 5 mg Nasal decongestant

Relieves:

Sinus congestion & pressure; Headache & pain; Runny nose & sneezing; Cough

Actual Size

8 SOFTGELS 20 TOTAL

NON DROWSY DAYTIME AND NIGHTTIME SINUS CONGESTION AND COUGH RELIEF

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride kit

Product Informat	ion			
Product Type	HUMAN OTC DRUG	Item Cod	e (Source)	NDC:69842-824
Packaging				
# Hom Codo	Dackaga Dacarintia	` `	Marketing Start	Marketing End

# item code	Package Description		Date	Da	ate
1 NDC:69842-824- 20 Product	CARTON; Type 0: Not a Combina 	ation 08/25/20	21		
Quantity of Parts					
Part # Pack	age Quantity	Tot	al Product Qu	uantity	
Part 1 1 BLISTER PACK	1	12			
Part 2 1 BLISTER PACK	8	3			
Part 1 of 2					
	AYTIME SINUS CO methorphan hydrobromid				
Product Information	n				
ltem Code (Source)	NDC:51316-891				
Route of Administratio	n ORAL				
Active Ingredient/Ac	t ive Moiety ngredient Name		Basis of St	renath	Strengt
	209ITL9D) (ACETAMINOPHEN - I	UNII: 36209ITI 9D)	ACETAMINOPHEN	-	325 mg
	DROBROMIDE (UNII: 9D2RTI9K		DEXTROMETHORF HYDROBROMIDE		10 mg
PHENYLEPHRINE HYDROC JNII:1WS297W6MV)	HLORIDE (UNII: 04JA59TNSJ) (P	HENYLEPHRINE -	PHENYLEPHRINE		5 mg
Inactive Ingredients	Ingredient Name			Str	ength
FD&C RED NO. 40 (UNII: W	-			Ju	engen
FD&C YELLOW NO. 6 (UNII					
GELATIN (UNII: 2G86QN327I					
GLYCERIN (UNII: PDC6A3C0					
POLYETHYLENE GLYCOL 4	00 (UNII: B697894SGQ)				
POVIDONE (UNII: FZ989GH9	94E)				
PROPYLENE GLYCOL (UNII:	6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)					
SORBITOL (UNII: 506T60A2					
SORBITAN (UNII: 6092ICV9F	RU)				
TITANIUM DIOXIDE (UNII: 1	5FIX9V2JP)				
Product Characteris	+:				

Color r	ed (Orange to red)	Score	no score
Shape C	OVAL (oblong)	Size	20mm
Flavor		Imprint Code	512;A09
Contains			

Pa	ckaging			
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1		1 in 1 CARTON		
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
· · ·				

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	M012	08/25/2021	

Part 2 of 2

NIGHTTIME SINUS CONGESTION AND COUGH RELIEF

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate, phenylephrine hydrochloride capsule, liquid filled

Product Information

Item Code (Source)	NDC:51316-892
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	5 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)	

POLYEIHYLER	NE GLYC	OL 400 (UNII: B697894SGQ)			
POVIDONE (UI					
		(UNII: 6DC9Q167V3)			
WATER (UNII: (059QF0K	O0R)			
SORBITOL (UN	III: 506T6	50A25R)			
SORBITAN (UN	III: 6092	ICV9RU)			
SODIUM HYDP	ROXIDE	(UNII: 55X04QC32I)			
TITANIUM DIO	XIDE (U	NII: 15FIX9V2JP)			
Product Cl	naract	eristics			
Color		green (transparent)	Sc	ore	no score
Shape		OVAL (oblong)	Si	ze	20mm
Flavor			Im	print Code	116;A07
Contains					
Packaging					
ltom		Package Description	1	Marketing Start Date	Marketing End Date
# Item	1 in 1 C		1		-
# Item Code		CARTON BLISTER PACK; Type 0: Not a Col			-
# Item Code 1	8 in 1 E	CARTON BLISTER PACK; Type 0: Not a Col			-
# Item Code 1	8 in 1 E Product	CARTON BLISTER PACK; Type 0: Not a Col			-
# Item Code 1	8 in 1 E Product	CARTON BLISTER PACK; Type 0: Not a Con	mbination		-
# Item Code 1 1 Marketir Categor	8 in 1 E Product	CARTON BLISTER PACK; Type 0: Not a Cont formation Application Number or M Citation	mbination	Date Date Marketing Start	Date Marketing End
# Item Code 1 1 Marketir Categor	8 in 1 E Product	CARTON BLISTER PACK; Type 0: Not a Cont formation Application Number or M Citation	mbination	Date Date	Date Marketing End
# Item Code 1 1 1 Marketir Categor OTC Monograp	8 in 1 E Product	CARTON BLISTER PACK; Type 0: Not a Cont formation Application Number or M Citation	mbination	Date Date	Date Marketing End
# Item Code 1 1 1 Marketir Categor OTC Monograp	8 in 1 E Product	CARTON BLISTER PACK; Type 0: Not a Cont formation Application Number or N Citation	mbination Monograph	Date Date	Date Marketing End

Labeler - CVS PHARMACY, INC. (062312574)

Revised: 12/2023

CVS PHARMACY, INC.