

D TIME SINUS- acetaminophen and phenylephrine hydrochloride capsule, liquid filled
SPIRIT PHARMACEUTICALS,LLC

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.

D-Time Sinus Soft Gelatin Capsules

Drug Facts

<i>Active ingredients for DayQuil Sinus (in each LiquiCap)</i>	<i>Purpose</i>
Acetaminophen 325 mg	Pain reliever
Phenylephrine HCl 5 mg	Nasal decongestant

Uses

temporarily relieves nasal and sinus symptoms:

- sinus pain
- headache
- nasal and sinus congestion
- runny nose and sneezing (NyQuil Sinus only)

Warnings

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

- more than 6 doses in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). Ask a doctor or pharmacist before using with other drugs if you are not sure
- 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.
- to make a child sleep

Ask a doctor before use if you have

- liver disease
- heart disease
- thyroid disease
- diabetes
- high blood pressure
- trouble urinating due to enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis
- glaucoma

Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers.

When using these products

- **do not use more than directed**

In addition, when using NyQuil Sinus:

- excitability may occur, especially in children
- marked drowsiness may occur
- avoid alcoholic drinks
- be careful when driving a motor vehicle or operating machinery
- alcohol, sedatives, and tranquilizers may increase drowsiness

Stop use and ask a doctor if

- redness or swelling is present
- symptoms do not get better within 7 days or are accompanied by a fever
- you get nervous, dizzy or sleepless
- fever gets worse or lasts more than 3 days
- new symptoms occur

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 health problems. 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.

Directions

- take only as recommended - see Overdose warning
- do not exceed 6 doses per 24 hours

NyQuil Sinus OR DayQuil Sinus

adults and children 12 years and over	2 LiquiCaps with water every 4 hours
children 2 to under 12 years	ask a doctor
children under 2 years	do not use

- **when using other DayQuil or NyQuil products, carefully read each label to insure correct dosing**

Other information

- store at room temperature

Inactive ingredients

NyQuil Sinus FD&C Blue No. 1, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol special, titanium dioxide.

DayQuil Sinus FD&C Yellow No. 6, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, sorbitol special, titanium dioxide.

PRINCIPAL DISPLAY PANEL - 6000 Softgels

D-Time Sinus Soft Gelatin Capsules

Each Softgel Contains:

Acetaminophen USP 325mg

Phenylephrine HCL USP 5mg

LOT NO :	NDC NO : 68210-1470-6
MFG DATE	QUANTITY : 6000 Softgels
:	
EXP. DATE	GROSS WT. :
:	

WARNING:

KEEP OUT OF REACH OF CHILDREN

STORE AT CONTROLLED ROOM TEMPERATURE OF 59° - 86°F (15° - 30°C)

PROTECT FROM LIGHT, MOISTURE AND FREEZING

**THIS BULK SHIPMENT IS INTENDED FOR FURTHER PACKAGING PROCESS ONLY.
CONTENTS SHOULD BE APPROVED, REPACKAGED IMMEDIATELY AND LABELED IN
STRICT**

CONFORMANCE WITH THE FD & C ACT AND REGULATIONS THEREUNDER.

MANUFACTURED BY:

Marksans Pharma Ltd

VERNA, GOA-403722,

INDIA.

CODE : GO/DRUGS/515

MANUFACTURED FOR:

SPIRIT PHARMACEUTICALS LLC

225 LINCOLN HWY, STE 205

FAIRLESS HILLS, PA 19030

PH.# 215 943 4000

FAX.# 215 943 4039

CAUTION : "FOR MANUFACTURING, PROCESSING OR REPACKING"

D-Time Sinus Soft Gelatin Capsules Each Softgel Contains: Acetaminophen USP 325mg Phenylephrine HCL USP 5mg	
LOT NO :	NDC NO : 68210-1470-6
MFG DATE :	QUANTITY : 6000 Softgels
EXP DATE :	GROSS WT. :
WARNING: KEEP OUT OF REACH OF CHILDREN	
STORE AT CONTROLLED ROOM TEMPERATURE OF 59° - 86°F (15° - 30°C) PROTECT FROM LIGHT, MOISTURE AND FREEZING	
THIS BULK SHIPMENT IS INTENDED FOR FURTHER PACKAGING PROCESS ONLY. CONTENTS SHOULD BE APPROVED, REPACKAGED IMMEDIATELY AND LABELED IN STRICT CONFORMANCE WITH THE FD & C ACT AND REGULATIONS THEREUNDER.	
MANUFACTURED BY: Marksans Pharma Ltd VERNA, GOA-403722, INDIA. CODE : GO/DRUGS/515	MANUFACTURED FOR: SPIRIT PHARMACEUTICALS LLC 225 LINCOLN HWY, STE 205 FAIRLESS HILLS, PA 19030 PH.# 215 943 4000 FAX.# 215 943 4039
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D TIME SINUS

acetaminophen and phenylephrine hydrochloride capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68210-1470
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GELATIN (UNII: 2G86QN327L)	
POVIDONE (UNII: FZ989GH94E)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Product Characteristics

Color	ORANGE	Score	no score
Shape	OVAL	Size	18mm
Flavor		Imprint Code	129
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68210-1470-5	1 in 1 DRUM		
1		5000 in 1 BAG		
2	NDC:68210-1470-6	1 in 1 DRUM		
2		6000 in 1 BAG		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part341	03/01/2010	

Labeler - SPIRIT PHARMACEUTICALS,LLC (179621011)

Revised: 7/2010

SPIRIT PHARMACEUTICALS,LLC