

**AXIM DAYTIME- acetaminophen, dextromethorphan hydrobromide,
phenylephrine hydrochloride capsule, liquid filled
VIVUNT PHARMA 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.

AXIM - DayTime Cold & Flu

Drug Facts

Active ingredients (in each softgel)	Purpose
Acetaminophen 325 mg	Pain reliever/fever reducer
Dextromethorphan HBr 10 mg	Cough suppressant
Phenylephrine HCl 5 mg	Nasal decongestant

Uses

Temporarily relieves common cold/flu symptoms:

- nasal Congestion
- cough due to minor throat and bronchial irritation
- sore throat
- headache
- minor aches and pains
- fever

Warnings

Liver warning

This product contains 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 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, 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

Ask a doctor before use if you have

- liver disease
- heart disease
- high blood pressure
- diabetes
- thyroid disease
- trouble urinating due to an 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 exceed recommended dose

Stop use and ask a doctor if

- you get nervous, dizzy or sleepless
- 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 or headache that lasts.

These could be signs of a serious condition

Keep out of reach of children.

If pregnant or breast-feeding

ask a health professional before use.

Overdose warning

In case of accidental overdose, seek professional assistance or contact a poison control

center immediately.

Directions

- Take only as directed (see overdose warning)
- Do not take more than 8 softgels in 24 hours.

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

Other information

- Store at 20° - 25 °C (68 °- 77 °F)
- Read all product information before using
- Tamper evident: Do not use if carton is open or blister unit is broken.

Inactive ingredients

FD&C Red No. 40, FD&C Yellow No. 6, Gelatin, Glycerin, Methylparaben, Polyethylene Glycol 400, Povidone K30, Propylene Glycol, Propylparaben, Purified Water, Sorbitol, Titanium Dioxide

Product of India

Distributed by:

VIVUNT PHARMA LLC
8950 SW 74th. Court. Suite 1901
Miami, Florida. Z,C. 33156-3178

PRINCIPAL DISPLAY PANEL - Axim DayTime Carton 24

Compare to

VICKS ® DayQuil®

Cold&Flu LiquiCaps®

active ingredients*

NDC 82706-001-01

AXIM - DayTime

COLD&FLU

Multi-Symptom Relief

- Pain Reliever
- Fever Reducer
- Cough Suppressant

- Nasal Decongestant

Non-Drowsy

Acetaminophen , Phenylephrine HCl,

Dextromethorphan HBr

24 SOFTGELS** **Liquid-filled capsules

*This product is not manufactured or distributed by

The Procter & Gamble Company, owner of the registered trademarks Vicks® DayQuil® and LiquiCaps®.



PRINCIPAL DISPLAY PANEL - Axim DayTime Carton 6

Compare to

VICKS® DayQuil®

Cold&Flu LiquiCaps®

active ingredients*

NDC 82706-001-02

AXIM - DayTime

COLD&FLU

Multi-Symptom Relief

Pain Reliever

Fever Reducer

Cough Suppressant

Nasal Decongestant

Non-Drowsy

Acetaminophen , Phenylephrine HCl,

Dextromethorphan HBr

6 SOFTGELS** **Liquid-filled capsules

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Miami, Florida Z.C. 33156-3178

AXIM DAYTIME				
acetaminophen, dextromethorphan hydrobromide, phenylephrine hydrochloride capsule, liquid filled				
Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:82706-001
Route of Administration		ORAL		
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)			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
METHYLPARABEN (UNII: A2I8C7HI9T)				
GLYCERIN (UNII: PDC6A3C0OX)				
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
GELATIN (UNII: 2G86QN327L)				
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)				
POVIDONE K30 (UNII: U725QWY32X)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
SORBITOL (UNII: 506T60A25R)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
Product Characteristics				
Color	orange	Score	no score	
Shape	OVAL	Size	22mm	
Flavor		Imprint Code	axim	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82706-001-01	3 in 1 CARTON	05/09/2022	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
	NDC 82706-001			

2	NDC:82706-001-02	3 in 1 CARTON	09/20/2023	
2		2 in 1 POUCH; 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		05/09/2022	

Labeler - VIVUNT PHARMA LLC (045829437)

Revised: 9/2023

VIVUNT PHARMA LLC