AQUANAZ- dextromethorphan hydrobromide, guaifenesin and phenylephrine hydrochloride tablet Capital Pharmaceutical, LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Aquanaz Tablets

Aquanaz Drug Facts.

<u>Active Ingredient.</u>

Dextromethorphan HBr 15mg.

Purpose.

Cough Suppressant.

<u>Active Ingredient.</u>

Guaifenesin 400mg.

Purpose.

Expectorant.

Active Ingredient.

Phenylephrine HCL 10mg.

<u>Purpose.</u>

Nasal Decongestant.

Uses:

Temporally relieves these symptoms accruing with a cold nasal decongestion cough due to minor throat and bronchial irritation—helps loosen phlegm (mucus) and thins bronchial secretions to drain bronchial tubes.

Warnings: When using this product do not exceed recommended dose.

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 MAOI drugs. If you do not know if your prescription drug contains an MAOI. Consult a doctor or pharmacist before taking this product.

Stop use and ask a doctor if:

Symptoms do not improve--new symptoms occur--redness or swelling is present-nervousness, dizziness or sleeplessness occurs--symptoms do not improve within 7 days or are accompanied by fever--cough persists for more than 1 week, tends to recur, or is a accompanied by fever, rash or persistent headache. A persistent cough may be the sign of a serious condition.

Ask a doctor before use if you have:

Heart disease--high blood pressure--thyroid disease—diabetes--difficulty in urination due to enlargement of the prostate gland—persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis or emphysema or where cough is accompanied by excessive phlegm (mucus)

Directions:

Adults and children 12 years and older: Take 1 tablet every 4-6 hours as needed, do not exceed 4 tablets in 24hours, or as directed by a doctor.

<u>Children 6-12 years:</u> Take $\frac{1}{2}$ tablet every 4-6 hours as needed, do not exceed 2 tablets in 24 hours, or as directed by a doctor.

If pregnant or breast-feeding baby,

ask a health care professional before use.

KEEP OUT OF REACH OF CHILDREN.

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.

Inactive Ingredients:

F&C Blue #2 Microcrystalline Celulose, Silicone Dioxide, Stearic Acid.

Other Information:

Do not use if there are signs of tampering—Store at controlled room temperature 15deg-30degC (59-86degF)

PRINCIPAL DISPLAY PANEL

NDC 29978-587-01 **NEW**

Aquanaz TABLETS

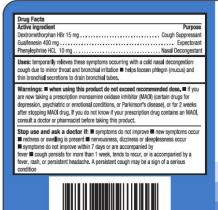
TRIPLE COMBINATION RELIEF:

Cough Suppressant: Dextromethorphan HBR 15 MG

Expectorant: Guaifenesin 400 MG

Decongestant: Phenylephrine HCL 10 MG

100 Tablets





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Drug Facts (Continued)

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Capital Pharmaceuticals, LLC. Powell, OH 43065

AQUANAZ

dextromethorphan hydrobromide, quaifenesin and phenylephrine hydrochloride tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:29978-587

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength **DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) DEXTROMETHORPHAN** 15 mg (DEXTROMETHORPHAN - UNII:7355X3ROTS) **HYDROBROMIDE** GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) 400 mg **GUAIFENES IN** PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE -**PHENYLEPHRINE** 10 mg UNII:1WS297W6MV) **HYDROCHLORIDE**

Inactive Ingredients			
Ingredient Name	Strength		
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)			
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)			
SILICON DIOXIDE (UNII: ETJ7Z 6XBU4)			
STEARIC ACID (UNII: 4ELV7Z65AP)			

Product Characteristics				
Color	blue (light blue)	Score	no score	
Shape	OVAL	Size	15mm	
Flavor		Imprint Code	PAC3	
Contains				

l	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:29978-587- 01	100 in 1 BOTTLE; Type 0: Not a Combination Product	03/10/2015	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/10/2015	

Labeler - Capital Pharmaceutical, LLC (831545541)

Revised: 2/2024 Capital Pharmaceutical, LLC