ANTIGRIP COUGH AND COLD- acetaminophen, chlorpheniramine maleate, dextromethorphan hbr, phenylephrine hcl tablet, effervescent
Pharmadel LLC

Antigrip Cough and Cold Purple Effervescent (V)

Drug Facts

Active ingredients & Purposes

Active ingredients (in each tablet)	***Purposes***
Acetaminophen 325 mg.....	**Pain reliever/ fever reducer**
Chlorpheniramine maleate 2 mg	Antihistamine
Dextromethorphan HBr 10 mg	Cough suppressant
Phenylephrine HCl 5 mg	Nasal decongestant

Uses

For the temporary relief of symptoms due to the common cold/ flu:

- runny nose
- sneezing
- nasal congestion
- cough due to minor throat and bronchial irritation
- minor aches and pain
- headache
- sore throat
- and temporarily reduces fever

Warnings

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

- more than **12 tablets 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

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, persists for more than 2 days, and 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 your 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
- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the prostate gland
- a persistent or chronic cough such as occurs with smoking, asthma, or emphysema
- a cough is accompanied by excessive phlegm (mucus)

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 the recommended dosage**
- may cause marked drowsiness
- may cause excitability, especially in children
- may cause drowsiness; alcohol, sedatives, and tranquilizers may increase the drowsiness effect
- avoid alcoholic beverages while taking this product
- use caution when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occurs
- 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 a rash or headache that lasts. These could be signs of a serious condition.

If pregnant or breastfeeding,

ask a health professional before use.

Keep out of reach of children.

In case of an accidental overdose, get medical help or contact a Poison Control Center

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

Directions

■ **adults and children 12 years of age and older:** take two (2) tablets fully dissolved in a 4 oz glass of water every 4 hours. Do not exceed **12 tablets in 24 hours,** or as directed by a doctor.

■ **children under 12 years of age:** do not use

Other information

- each tablet contains: **potassium 10mg, sodium 290mg**
- store below 77° F (25° C) and in a dry place

Inactive ingredients

acesulfame potassium, anhydrous citric acid, FD&C blue #1, FD&C red #40, grape flavor, polysorbate 80, povidone k-30, simethicone, sodium benzoate, sodium bicarbonate, sucralose, water

Questions or comments?

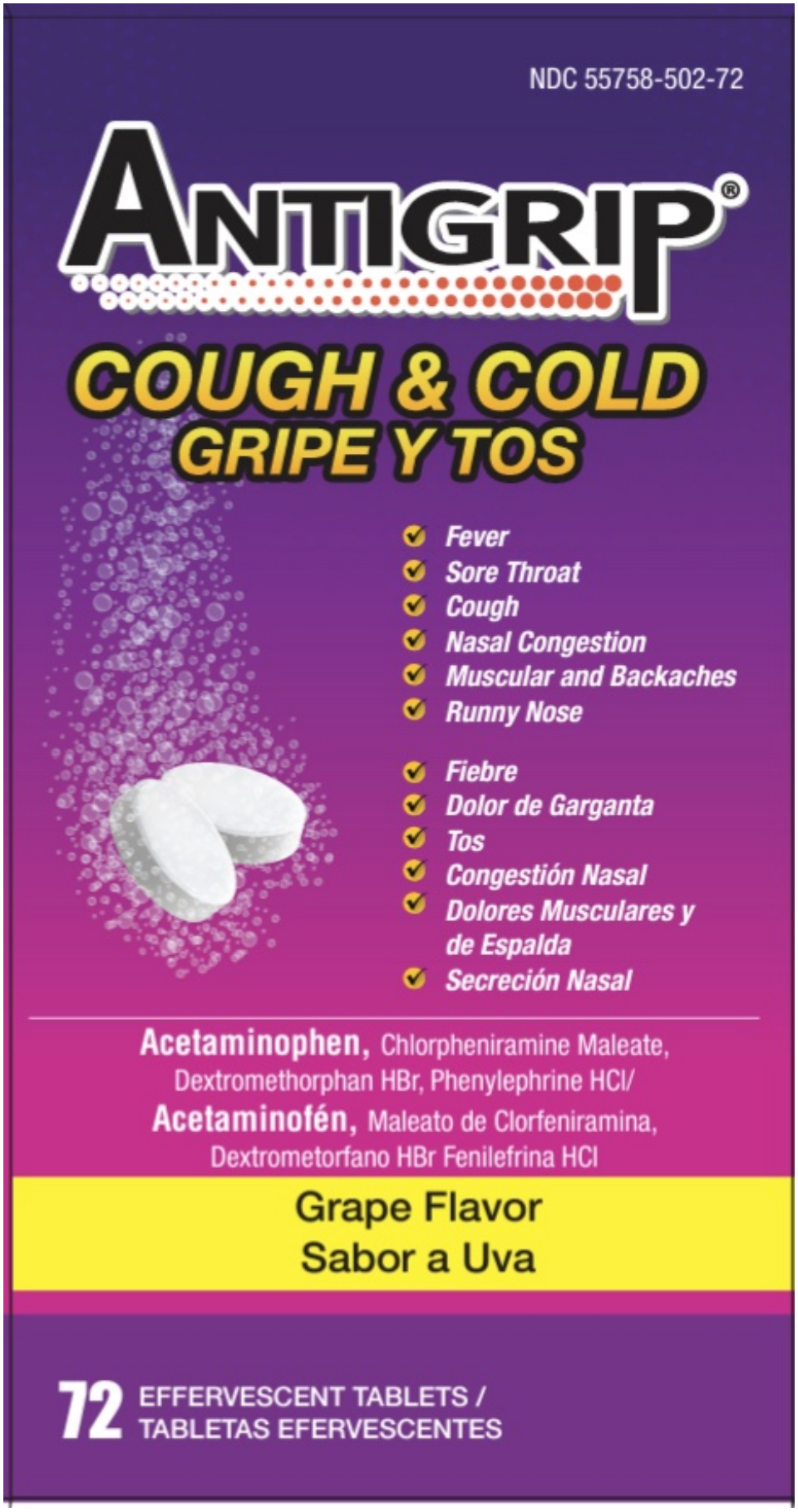
+1-866-359-3478 (M-F) 9 AM to 5 PM EST or www.pharmadel.com

Dist. by:

Pharmadel LLC.

New Castle, DE 19720

PRINCIPAL DISPLAY PANEL



ANTIGRIP COUGH AND COLD

acetaminophen, chlorpheniramine maleate, dextromethorphan hbr, phenylephrine hcl tablet,

effervescent

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55758-502
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
DIMETHICONE (UNII: 92RU3N3Y1O)	

Product Characteristics

Color	gray	**Score**	no score
Shape	ROUND	**Size**	24mm
Flavor	GRAPE	**Imprint Code**	V3
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55758-502-72	36 in 1 CARTON	11/01/2024	
1		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 Drug	M012	11/01/2024	

Labeler - Pharmadel LLC (030129680)

Revised: 10/2024 Pharmadel LLC