**ANTIGRIP NIGHTTIME- acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl powder, for solution**
**Pharmadel LLC**

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**Antigrip Nighttime Tea G-H-L (AS)**

**Drugs Facts**

**Active Ingredients & Purposes**

| ***Active ingredients (in each packet)*** | ***Purposes*** |
|---|---|
| **Acetaminophen 325 mg.....** | **Pain reliever/ fever reducer** |
| Dextromethorphan HBr 10 mg..... | Cough suppressant |
| Doxylamine succinate 6.25 mg ..... | Antihistamine |
| Phenylephrine HCl 5 mg..... | Nasal decongestant |

***Uses***

Temporary relieves common cold/flu symptoms:

- minor aches and pains
- sore throat
- headache
- nasal congestion
- runny nose

- itching of nose or throat
- stuffy nose
- cough due to minor throat and bronchial irritation
- and temporarily reduces fever

***Warnings***

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

- more than 6 packets 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.

**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 prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or 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

**Ask a doctor before use if you**

- have liver disease
- have heart disease
- have high blood pressure
- have thyroid disease
- have diabetes
- have glaucoma
- have a cough that is accompanied by excessive phlegm (mucus)
- have a persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- have difficulty urinating due to an enlarged prostate gland
- are taking sedatives or tranquilizers

**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**
- may cause excitability, especially in children
- use caution when driving a motor vehicle or operating machinery
- may cause marked drowsiness

**Stop use and ask doctor if**

- nervousness, dizziness, or sleeplessness occurs
- redness or swelling is present
- pain, cough, or nasal congestion gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- any new symptoms occur
- cough comes back or occurs with rash or headache that lasts.

These could be signs of a serious condition.

**If pregnant or breast-feeding,**

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***

- dissolve one packet in an 8oz. glass of hot water. If using a microwave; add content into a glass of cool water and stir briskly before and after heating. Do not overheat.
- sip while hot and consume the entire drink within 10-15 minutes

| Age | Dose |
|---|---|
| adults and children 12 years of age and over | one (1) packet every 4 hours,<br>take every 4 hours; do not exceed 6 packets in a 24 hours |
| children under 12 years of age | do not use |

***Other information***

- each packet contains: **potassium 10 mg, sodium 27 mg**
- **phenylketonurics:** contains phenylalanine 13 mg per packet
- store at room temperature 68-77°F (20-25°C)
- do not use if packet is punctured or torn

***Inactive ingredients***

acesulfame potassium, anhydrous citric acid, aspartame, isopropyl alcohol, lemon ginger honey flavors, maltodextrin, silicon dioxide, sodium citrate, sucrose, tribasic calcium phosphate, water

**Questions & 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 NIGHTTIME

acetaminophen, dextromethorphan hbr, doxylamine succinate, phenylephrine hcl powder, for solution

## Product Information

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:55758-382 |
| **Route of Administration** | ORAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **DOXYLAMINE SUCCINATE** (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL) | DOXYLAMINE SUCCINATE | 6.25 mg |
| **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 |
|---|---|
| **ANHYDROUS CITRIC ACID** (UNII: XF417D3PSL) | |
| **ISOPROPYL ALCOHOL** (UNII: ND2M416302) | |
| **MALTODEXTRIN** (UNII: 7CVR7L4A2D) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **SILICON DIOXIDE** (UNII: ETJ7Z6XBU4) | |
| **ACESULFAME POTASSIUM** (UNII: 23OV73Q5G9) | |

| | |
|---|---|
| **ASPARTAME** (UNII: Z0H242BBR1) | |
| **SODIUM CITRATE** (UNII: 1Q73Q2JULR) | |
| **SUCROSE** (UNII: C151H8M554) | |
| **TRIBASIC CALCIUM PHOSPHATE** (UNII: 91D9GV0Z28) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | | **Score** | |
| **Shape** | | **Size** | |
| **Flavor** | LEMON (Lemon - Ginger - Honey) | **Imprint Code** | |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:55758-382-18 | 18 in 1 CARTON | 03/28/2024 | |
| **1** | NDC:55758-382-01 | 1 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 | 03/28/2024 | |

## Labeler - Pharmadel LLC (030129680)

Revised: 3/2024 Pharmadel LLC