ANTIGRIP- acetaminophen, phenylephrine hcl, chlorpheniramine maleate, dextromethorphan hbr capsule, liquid filled HEALTHLIFE OF USA LLC

HealthLife [®]Antigrip Capsule

Active ingredients (in each softgel)

- Acetaminophen 250mg
- Chlorpheniramine maleate 2 mg
- Dextromethorphan hydrobromide 10 mg
- Phenylephrine hydrochloride 5 mg

Purpose

Pain reliever/fever reducer

Antihistamine

Cough suppressant

Nasal decongestant

Uses

- temporarily relieves these symptoms due to a cold or flu:
- minor aches and pains
- headache
- nasal and sinus congestion
- cough
- runny nose
- sneezing
- sore throat
- temporarily reduces fever

Warnings

Liver warning

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

- more than 10 softgels 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, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

Do not use to sedate children.

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
- in children under 12 years of age

Ask a doctor before use if you have

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- glaucoma
- cough with excessive phlegm (mucus)
- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema

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

Stop use and ask a doctor if

- 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 rash or headache that lasts.

These could be signs of a serious condition.

• nervousness, dizziness, or sleeplessness occurs

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. 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

- do not take more than the recommended dose
- adults and children 12 years and over: take 2 softgels with water every 4 hours. Do not exceed 10 softgels in 24 hours or as directed by a doctor.
- children under 12 years: do not use

Other information

• Store at room temperature 15°-30°C (59°F-86°F) and avoid excessive heat.

Inactive ingredients

FD&C blue no.1, FD&C red no. 33, gelatin, glycerin, polyethylene glycol, povidones,

propylene glycol, shellac, sodium hydroxide, sorbitan, sorbitol, titanium dioxide, water.

Questions or commonts

1-844-832-1138 Monday through Friday 9am-5pm

PRINCIPAL DISPLAY PANEL - Shipping Label

Cough and Cold plus Capsules Each softgel contains: Acetaminophen 250 mg Phenylephrine HCl 5 mg Chlorpheniramine Maleate 2 mg Dextromethorphan HBr 10 mg Quantity: 8 Softgels NDC No.:69517-112-08 IMPORTANT: 1. Inspect immediately upon receipt.

- 2. This is a bulk shipment, intended for further processing only.
- 3. Protect from heat, humidity, and light. Do not refrigerate.
- 4. 4.Store at 15-30°C (59-86)°F



ANTIGRIP

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Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:69517-112	
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
Ingredient Name			Basis of Str	ength	Strength
ACETAMINOPHEN (UNII: 36209IT	L9D) (ACETAMINOPHEN - UNII	:362O9ITL9D)	ACETAMINOPHEN		250 mg
PHENYLEPHRINE HYDROCHLOR UNII:1WS297W6MV)	IYLEPHRINE -	PHENYLEPHRINE HYDROCHLORIDE		5 mg	
CHLORPHENIRAMINE MALEATE UNII:3U6IO1965U)	(UNII: V1Q0O9OJ9Z) (CHLORF	PHENIRAMINE -	CHLORPHENIRAMII MALEATE	NE	2 mg

		Ingredient Nan	no		Strength
FD&C BLUE NO. 1		-			Strength
GELATIN (UNII: 2G					
GLYCERIN (UNII: P					
		ECIFIED (UNII: 3MQ0SI	DWIA)		
POVIDONE (UNII: F			,		
PROPYLENE GLYC		Q167V3)			
SORBITOL (UNII: 5	06T60A25R)				
WATER (UNII: 0590	QF0KO0R)				
TITANIUM DIOXID	E (UNII: 15FIX9)	/2JP)			
METHYLPARABEN	(UNII: A2I8C7H	I9T)			
PROPYLPARABEN					
Product Char	actorictica				
			C	-	
	Color blue		Score	8	no score
Shape CAPSULE (Oblong)		(Oblong)	Size		21mm
			B		
Flavor			Impri	int Code	SN;5
			Impri	nt Code	SN;5
			Impri	nt Code	SN;5
Contains			Impri	nt Code	5N;5
Contains					
Contains Packaging	Pa	ickage Descriptior		nt Code Marketing Start Date	SN;5 Marketing End Date
Contains Packaging			1	Marketing Start	Marketing End
Contains Packaging # Item Code 1 NDC:69517-112-	1 in 1 CARTON		n	Marketing Start Date	Marketing End
Contains Packaging # Item Code 1 NDC:69517-112- 08	1 in 1 CARTON 8 in 1 BLISTEF	1	n	Marketing Start Date	Marketing End
Contains Packaging # Item Code 1 NDC:69517-112- 08	1 in 1 CARTON 8 in 1 BLISTEF	1	n	Marketing Start Date	Marketing End
Contains Packaging # Item Code 1 NDC:69517-112- 08 1	1 in 1 CARTON 8 in 1 BLISTEF Product	I R PACK; Type 0: Not a Co	n	Marketing Start Date	Marketing End
Contains Packaging # Item Code 1 NDC:69517-112- 08	1 in 1 CARTON 8 in 1 BLISTER Product	I R PACK; Type 0: Not a Co	ombination	Marketing Start Date	Marketing End

Labeler - HEALTHLIFE OF USA LLC (079656178)

Revised: 10/2023

HEALTHLIFE OF USA LLC