

**GANMAOLING- acetaminophen and chlorpheniramine maleate tablet**  
**ANHUI DONGSHENGYOUBANG PHARMACEUTICAL CO., LTD.**

*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.*

-----

**DRUG FACTS**

***Active ingredient (in each tablet)***

Acetaminophen 60 mg

Chlorpheniramine maleate 0.667 mg

***Purpose***

Pain reliever-fever reducer

Antihistamine

***Uses***

Temporarily relieves the following symptoms associated with the common cold, hay fever, or other upper respiratory allergies:

minor aches and pains

headache

muscular aches

fever

sore throat

sneezing

itching of the nose or throat

runny nose

itchy, watery eyes

***Warnings***

**Do not use**

for pain for more than 10 days

for fever for more than 3 days

**Ask a doctor before use if the user has**

glaucoma

a breathing problem such as emphysema or chronic bronchitis

difficulty in urination due to enlargement of the prostate gland

**Ask a doctor or pharmacist before use if you are**

taking sedatives or tranquilizers

**When using this product**

may cause excitability especially in children

may cause drowsiness (alcohol, sedatives, and tranquilizers may increase the drowsiness effect)

avoid alcoholic beverages

use caution when driving a motor vehicle or operating machinery

**Stop use and ask a doctor if**

pain or fever persists or worsen

new symptoms occur

redness or swelling is present

sore throat:

is severe

last for more than 2 days

is accompanied or followed by:

fever

headache

rash

nausea

vomiting

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. 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 6 tablets every 4 hours, not more than 36 tablets in 24 hours

children 6 to under 12 years of age: Take 3 tablets every 4 hours, not more than 18 tablets in 24 hours

children under 6 years: consult a doctor

***Other information***

protect from light and excessive heat

keep tightly closed

keep in dry place

***Inactive ingredients***

Rough-leaved holly root, evodia root, wild chrysanthemum flower, Chinese chaste tree twig, honeysuckle flower, strobilanthes cusia root, cornstarch, sucrose, water, FD&C yellow no.5 and FD&C yellow no.6.

**Questions or Comments? (888) 221-3496 M-F 9 am to 5 pm**

you may also use this number to report serious adverse events associated with the use of this product

GANMAOLING TABLETS, NDC 72030-002-01, Antihistamine, Pain Reliever-Fever Reducer, 120



Tablets

## GANMAOLING

acetaminophen and chlorpheniramine maleate tablet

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72030-002
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	60 mg
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	0.667 mg

### Inactive Ingredients

Ingredient Name	Strength
ILEX ASPRELLA ROOT (UNII: S7K9P1V8VG)	
MELICOPE PTELEIFOLIA ROOT (UNII: Z400593S4G)	
CHRYSANTHEMUM INDICUM FLOWER (UNII: I6OER6U04L)	
VITEX NEGUNDO WHOLE (UNII: C92PGK11XB)	
LONICERA JAPONICA FLOWER (UNII: 4465L2WS4Y)	
STROBILANTHES CUSIA ROOT (UNII: F1919HP06B)	
STARCH, CORN (UNII: O8232NY3SJ)	
SUCROSE (UNII: C151H8M554)	
WATER (UNII: 059QF0K00R)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

### Product Characteristics

Color	yellow	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	GM
Contains			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72030-002-01	1 in 1 BOX	11/26/2018	
1		120 in 1 BOTTLE; 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	11/26/2018	

**Labeler** - ANHUI DONGSHENGYOUBANG PHARMACEUTICAL CO., LTD. (527929527)

### Establishment

Name	Address	ID/FEI	Business Operations
ANHUI DONGSHENGYOUBANG PHARMACEUTICAL CO., LTD.		527929527	manufacture(72030-002)

Revised: 11/2018

ANHUI DONGSHENGYOUBANG PHARMACEUTICAL CO., LTD.