

# **GAMMA -12- interleukin-12 human recombinant - interferon gamma-1b - solution/ drops**

**Guna spa**

*Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.*

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## **GAMMA 12**

### **ACTIVE INGREDIENTS/PURPOSE**

INTERFERON GAMMA 4C IMMUNE SUPPORT  
INTERLEUKIN-12 4C IMMUNE SUPPORT

### **USES**

For the temporary relief of upper respiratory symptoms due to hay fever or airborne allergies such as: -Cough -Airway irritation -Shortness of breath

### **WARNINGS**

Stop use and ask doctor if symptoms of cough, shortness of breath, or wheezing do not improve, worsen or persist more than 3 days

### **PREGNANCY**

If pregnant or breast-feeding ask a doctor before use

### **WARNINGS**

Keep this and all medicines out of reach of children

### **DIRECTIONS**

Take 15 minutes before meals

#### **Adults and children 12 years and older**

20 drops in a little water, 2 times per day

#### **Children between 12 years and 6 years of age**

10 drops in a little water, 2 times per day

#### **Children under 6 years**

5 drops in a glass of water, 2 times per day

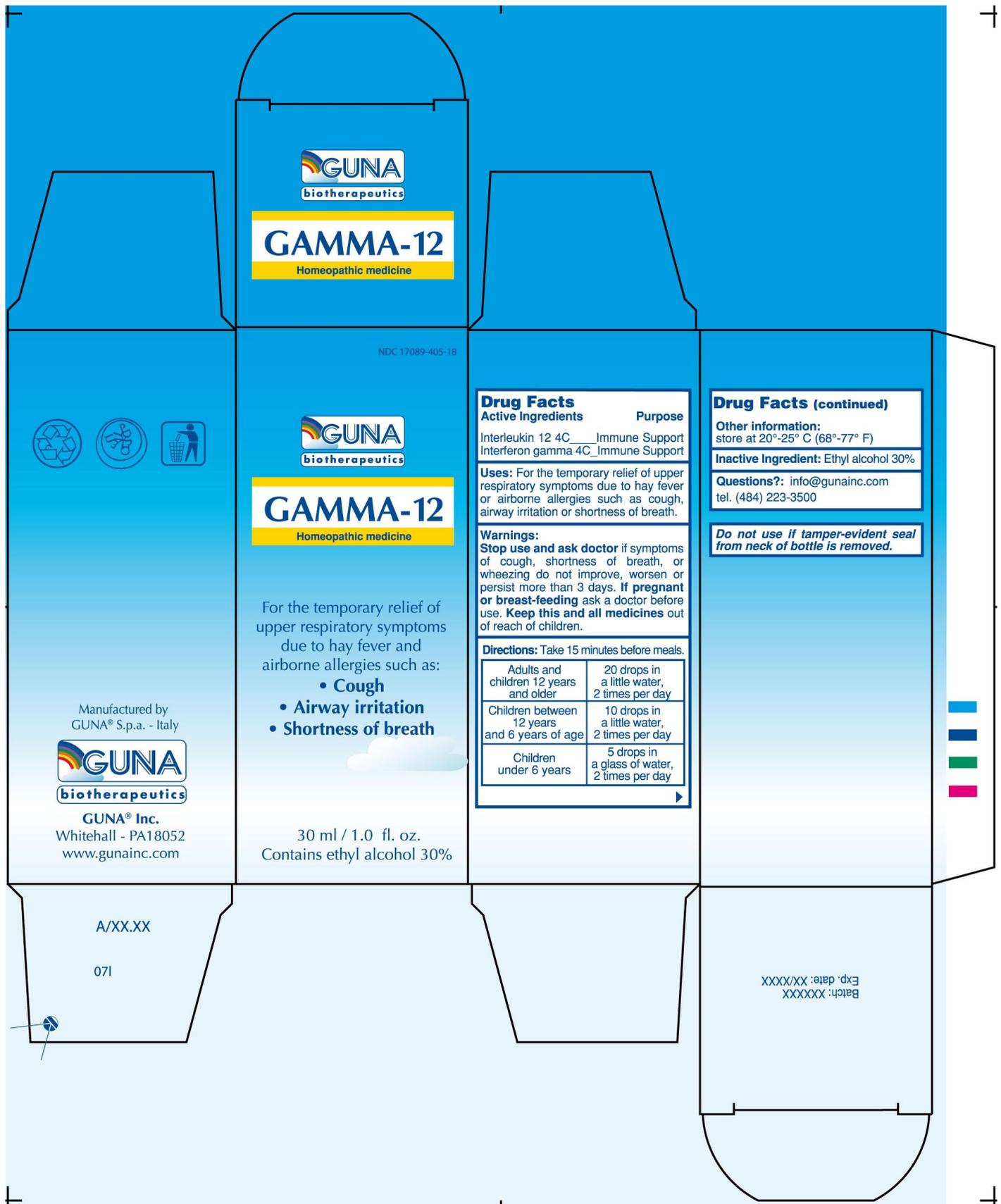
## **QUESTIONS**

Questions?: [info@gunainc.com](mailto:info@gunainc.com)  
tel. (484) 223-3500

inactive ingredient: Ethyl alcohol 30%.

take 15 minutes before meals

## **PRINCIPAL DISPLAY PANEL**



## GAMMA -12

interleukin-12 human recombinant - interferon gamma-1b - solution/ drops

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:17089-405
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>INTERFERON GAMMA-1B</b> (UNII: 21K6M2I7AG) (INTERFERON GAMMA-1B - UNII:21K6M2I7AG)	INTERFERON GAMMA-1B	4 [hp_C] in 30 mL
<b>INTERLEUKIN-12 HUMAN RECOMBINANT</b> (UNII: 02FXP10O2U) (INTERLEUKIN-12 HUMAN RECOMBINANT - UNII:02FXP10O2U)	INTERLEUKIN-12 HUMAN RECOMBINANT	4 [hp_C] in 30 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>ALCOHOL</b> (UNII: 3K9958V90M)	9 mL in 30 mL

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17089-405-18	1 in 1 BOX	03/31/2011	
1		30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		04/29/2010	

**Labeler** - Guna spa (430538264)

### Establishment

Name	Address	ID/FEI	Business Operations
Guna spa		338587646	manufacture(17089-405)

Revised: 11/2022

Guna spa