

GUNA-ANTI IL 1- anti-interleukin-1.alpha. immunoglobulin g rabbit- canakinumab 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.

GUNA-ANTI IL1

ACTIVE INGREDIENTS/PURPOSE

ANTI INTERLEUKIN 1 ALPHA	4C	ACUTE PAIN AND FEVER RELIEF
ANTI INTERLEUKIN 1 BETA	4C	ACUTE PAIN AND FEVER RELIEF

USES

Temporary relief of:

- **Acute pain**
- **Fever**

WARNINGS

- **Stop use and ask doctor if symptoms persist** more than 5 days.
- **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.
- Contains ethyl alcohol 30%

PREGNANCY

If pregnant or breast-feeding ask a doctor before use.

WARNINGS

Keepout of reach of children.

DIRECTIONS

Adults and children 12 years and older 20 drops twice a day in a little water. Hold in the mouth for about 30 seconds then swallow.

Children between 12 years and 6 years of age 10 drops twice a day in a little water. Hold in the mouth for about 30 seconds then swallow.

Children under 6 years 5 drops twice a day in a glass of water.

QUESTIONS

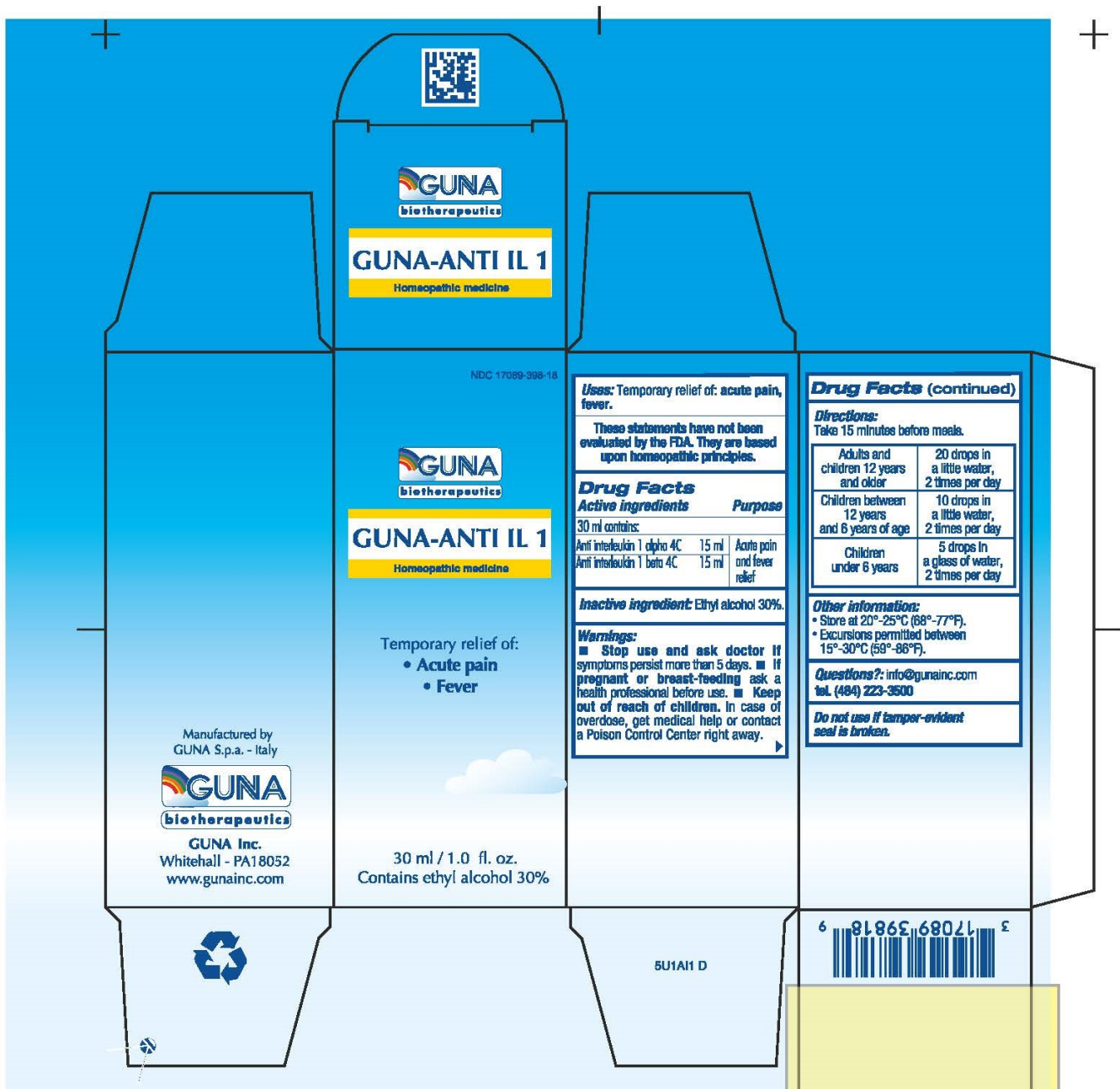
Questions?: info@gunainc.com

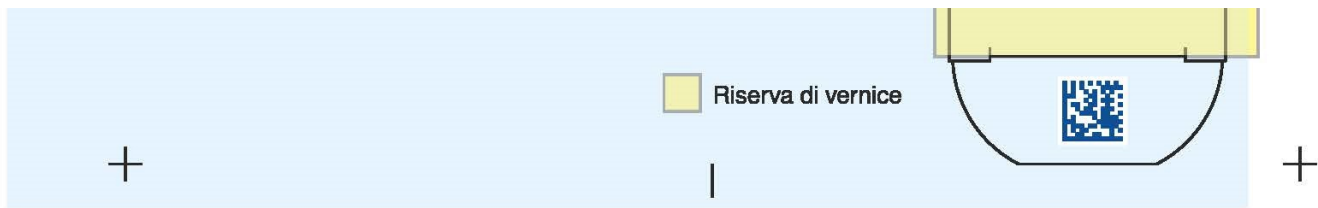
Tel. (484) 223-3500

Take 15 minutes before meals.

Inactive ingredient: Ethyl alcohol 30%.

PRINCIPAL DISPLAY PANEL





GUNA-ANTI IL 1

anti-interleukin-1.alpha. immunoglobulin g rabbit- canakinumab solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:17089-398
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ANTI-INTERLEUKIN-1.ALPHA. IMMUNOGLOBULIN G RABBIT (UNII: ML4QRZ1HCL) (ANTI-INTERLEUKIN-1.ALPHA. IMMUNOGLOBULIN G RABBIT - UNII:ML4QRZ1HCL)	ANTI-INTERLEUKIN-1.ALPHA. IMMUNOGLOBULIN G RABBIT	4 [hp_C] in 30 mL
CANAKINUMAB (UNII: 37CQ2C7X93) (CANAKINUMAB - UNII:37CQ2C7X93)	CANAKINUMAB	4 [hp_C] in 30 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	9 mL in 30 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17089-398-18	1 in 1 BOX	12/21/2018	
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		06/17/2008	

Labeler - Guna spa (430538264)

Establishment

Name	Address	ID/FEI	Business Operations
Guna spa		338587646	manufacture(17089-398)

Revised: 3/2020

Guna spa