

**GUNA-TF HERPES - human herpesvirus 1 - human herpesvirus 2 - capsule, gelatin coated
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.

DRUG FACTS

ACTIVE INGREDIENTS/PURPOSE

TRANSFER FACTOR HV1 7X IMMUNE STRENGTHENING
TRANSFER FACTOR HV2 7X IMMUNE STRENGTHENING

USES

For the temporary relief of symptoms due to cold sores: pain, tingling, burning sensation.

WARNINGS

Stop use and ask doctor if symptoms of tingling or burning sensation worsen or persist more than 5 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 1 capsule per day with a little water

Children between 12 years and 6 years of age 1 capsule every other day, with a little water

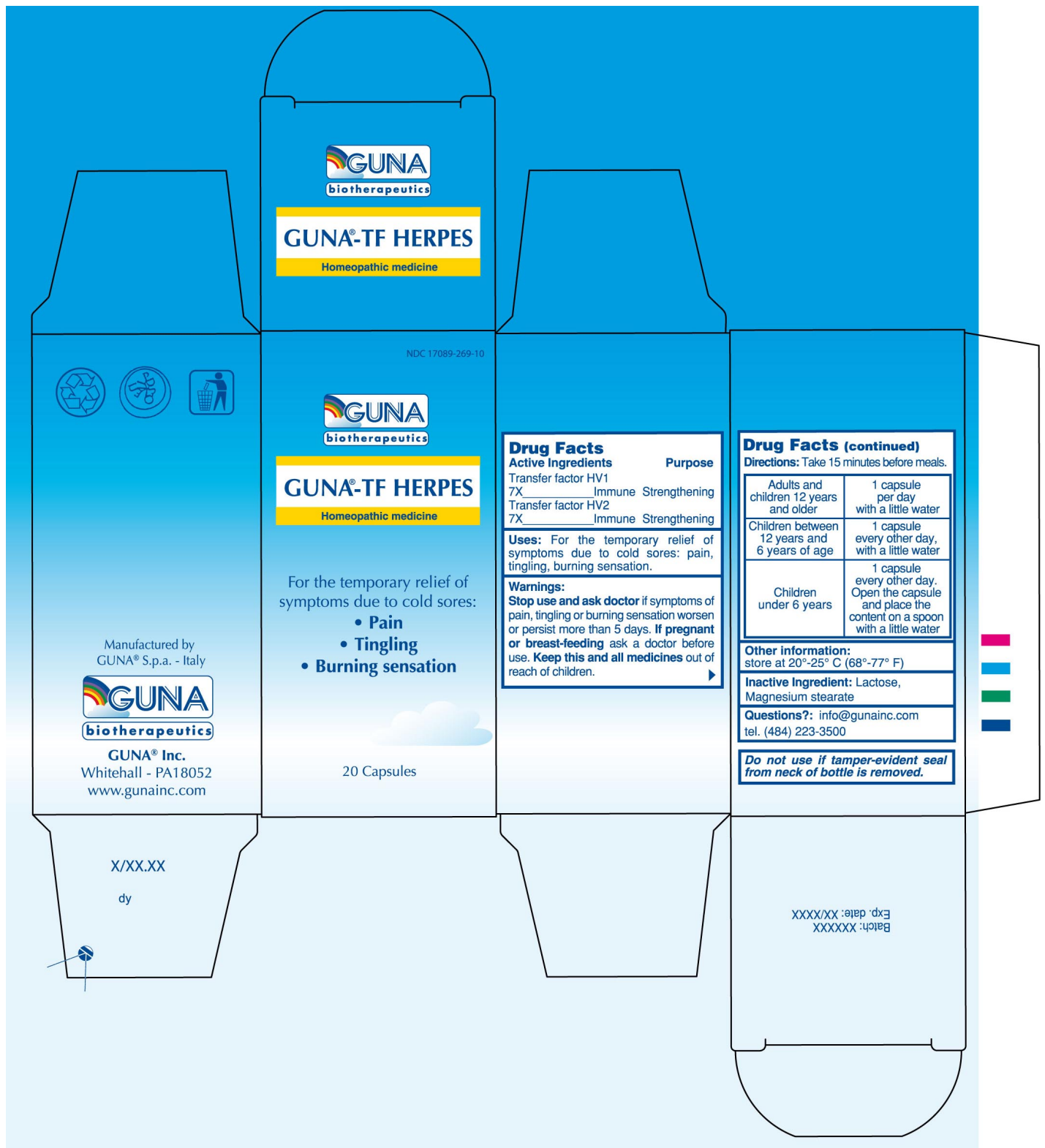
Children under 6 years 1 capsule every other day. Open the capsule and place the content on a spoon with a little water

QUESTIONS

Questions?: info@gunainc.com

Tel. (484) 223-3500

PRINCIPAL DISPLAY PANEL



GUNA-TF HERPES

human herpesvirus 1 - human herpesvirus 2 - capsule, gelatin coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:17089-269
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HUMAN HERPESVIRUS 1 (UNII: 22G38P19RL) (HUMAN HERPESVIRUS 1 - UNII:22G38P19RL)	HUMAN HERPESVIRUS 1	7 [hp_X] in 4600 mg
HUMAN HERPESVIRUS 2 (UNII: 74J6DNH49U) (HUMAN HERPESVIRUS 2 - UNII:74J6DNH49U)	HUMAN HERPESVIRUS 2	7 [hp_X] in 4600 mg

Inactive Ingredients

Ingredient Name	Strength
LACTOSE (UNII: J2B2A4N98G)	4048 mg in 4600 mg
MAGNESIUM STEARATE (UNII: 70097M6I30)	92 mg in 4600 mg

Product Characteristics

Color	white (white)	Score	2 pieces
Shape	OVAL (Capsule)	Size	18mm
Flavor		Imprint Code	na
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17089-269-10	1 in 1 BOX		
1		4600 mg in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		05/23/2006	

Labeler - Guna spa (430538264)**Establishment**

Name	Address	ID/FEI	Business Operations
Guna spa		430538264	manufacture

Revised: 3/2010

Guna spa