GEUM GENTIAN 1 SPECIAL ORDER- geum gentian 1 special order pellet Uriel Pharmacy Inc.

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.

Geum Gentian 1 Special Order

Directions: FOR ORAL USE ONLY.

Dissolve pellets under the tongue 3-4 times daily. Ages 12 and older: 10 pellets. Ages 2-11: 5 pellets. Under age 2: Consult a doctor.

Active Ingredients: 100 gm contains: 1 gm Gentiana e rad. 1X, 1 gm Geum urbanum e rad. 1X

Inactive Ingredient: Organic sucrose

Use: Temporary relief of digestive upset.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Contains sugar. Diabetics and persons intolerant of sucrose (sugar): Consult a doctor before use. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use. Do not use if safety seal is broken or missing.

Questions? Call 866.642.2858 Uriel, East Troy, WI 53120 www.urielpharmacy.com



GEUM GENTIAN 1 SPECIAL ORDER

geum gentian 1 special order pellet

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:48951-50	78
Route of Administration	ORAL				
Active Ingredient/Active M	Ioiety				
Active Ingredient/Active N	Aoiety Ingredient Name		Basis o	of Strength	Strength
Active Ingredient/Active M GENTIANA LUTEA ROOT (UNII: UNII:S7203284MS)	Ingredient Name	ROOT -	Basis o GENTIANA ROOT	0	Strength 1 [hp_Q]
GENTIANA LUTEA ROOT (UNII:	Ingredient Name S72O3284MS) (GENTIANA LUTEA		GENTIANA ROOT	0	U

Inactive Ingredients

Ingredient Name					Strength		
SUCROSE (UNII: C151H8 M554)							
Product Characteristics							
Color		white	Score		no score		
Shape		ROUND	Size		3mm		
Flavor			Imprint Code				
Contains							
Packaging							
# Item Code		Package Description		Marketing Star Date	rt Marketing End Date		
1 NDC:48951-5078- 2	1350 in 1 Product	50 in 1 BOTTLE, GLASS; Type 0: Not a Combination oduct		09/01/2009			
Marketing Information							
Marketing Catego	ory Aj	pplication Number or M	Ionograph Citation	Marketing Start Da	te Marketing End Date		
		09/01/2009					

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment						
Name	Address	ID/FEI	Business Operations			
Uriel Pharmacy Inc.		043471163	manufacture(48951-5078)			

Revised: 6/2018

Uriel Pharmacy Inc.