GENTIAN GINGER BITTERS- gentian ginger bitters liquid
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.

Gentian Ginger Bitters

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily before or after meals and/or snacks. Ages 12 and older: 1/4 teaspoon. Ages 2-11: 1/8 teaspoon. Under age 2: Consult a doctor.

Active Ingredients: 1gm contains: 350mg Gentiana (Yellow gentian) 1X, 277mg Absinthium (Wormwood) 1X, 223mg Zingiberis (Ginger root) 1X, 36.7mg Calamus (Sweet flag) 1X, 6.7mg Piper nigrum (Black peppercorns) 1X

Inactive Ingredient: Distilled water

Uses: Improves digestion; eases gas and indigestion.

KEEP OUT OF REACH OF CHILDREN.

Warnings: 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
Made with care by Uriel, East Troy, WI 53120
www.shopuriel.com Lot:

GENTIAN GINGER BITTERS
gentian ginger bitters liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-5006
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACORUS CALAMUS ROOT (UNII: XY1K7KIQ0F) (ACORUS CALAMUS ROOT - UNII:XY1K7KIQ0F)	ACORUS CALAMUS ROOT	1 [hp_X] in 1 mL
WHITE PEPPER (UNII: M29DW54Q9E) (WHITE PEPPER - UNII:M29DW54Q9E)	WHITE PEPPER	1 [hp_X] in 1 mL
WORMWOOD (UNII: F84709P2XV) (WORMWOOD - UNII:F84709P2XV)	WORMWOOD	1 [hp_X] in 1 mL
GINGER (UNII: C5529G5JPQ) (GINGER - UNII:C5529G5JPQ)	GINGER	1 [hp_X] in 1 mL
GENTIANA LUTEA ROOT (UNII: S72O3284MS) (GENTIANA LUTEA ROOT - UNII:S72O3284MS)	GENTIANA LUTEA ROOT	1 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-5006-3	125 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	09/01/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2009	

Labeler - Uriel Pharmacy, Inc. (043471163)

Establishment

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy, Inc.		043471163	manufacture(48951-5006)

Revised: 5/2022 Uriel Pharmacy, Inc.