GAS RELIEF ULTRA STRENGTH- simethicone capsule, liquid filled
Preferred Pharmaceuticals Inc.

Drug Facts

Active ingredient (in each softgel)

Simethicone 180 mg

Purpose

Antigas

Uses

- relieves bloating, pressure, or fullness commonly referred to as gas

Warnings

Stop use and ask a doctor if

condition persists.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

Directions

- swallow one or two softgels as symptoms occur
- do not exceed two softgels per 24 hours except under the advice and supervision of a physician

Other information

- store between 15-30ºC (59-86ºF)
- protect from heat and moisture

Inactive ingredients

FD&C red #40, FD&C yellow #6, gelatin, glycerin

Questions or comments?

Call **1-800-645-2158**

Principal Display Panel

Compare to the active ingredient in Phazyme® Ultra Strength†

Ultra Strength

Simethicone 180 mg

Antigas

Fast relief of

- Gas
- Pressure
- Bloating
- Discomfort

†This product is not manufactured or distributed by C.B. Fleet Company Inc., distributor of Phazyme® Ultra Strength

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Distributed by:

RUGBY® LABORATORIES

17177 N Laurel Park Drive,

Suite 233

Livonia, MI 48152

Error! Hyperlink reference not valid.

.

Product Label

RUGBY LABORATORIES Ultra Strength Simethicone

GAS RELIEF ULTRA STRENGTH

simethicone capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68788-7890(NDC:0536-1306)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	180 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

Product Characteristics

Color	ORANGE	**Score**	no score
Shape	CAPSULE	**Size**	12mm
Flavor		**Imprint Code**	05A
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-7890-3	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/23/2021	

2	NDC:68788-7890-6	36 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/23/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M002	04/23/2021	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-7890)

Revised: 8/2024 Preferred Pharmaceuticals Inc.