

**ULTRA STRENGTH GAS RELIEF- simethicone capsule, liquid filled
Wal-Mart Stores Inc**

Ultra Strength Gas Relief

Active ingredient (in each softgel)

Simethicone 180 mg

Purpose

Antigas

Use

for the relief of

- pressure and bloating commonly referred to as gas

Warnings

Keep out of reach of children

Directions

- adults: swallow with water 1 or 2 softgels as needed after meals and at bedtime
- do not exceed 2 softgels in 24 hours under the advice and supervision of a physician

Other information

- store at controlled room temperature 15° - 30°C (59° - 86°F)
- protect from heat and moisture
- do not use if imprinted safety seal under cap is broken or missing

Inactive ingredients

FD&C yellow #6, gelatin, glycerin, purified water and white ink

Questions or comments?

Call toll free: **1-888-287-1915**

PRINCIPAL DISPLAY PANEL

Ultra Strength Gas Relief

Simethicone 180 mg Anti-gas 50 Softgels

NDC 49035-585-21

*Compare to the active ingredient in Gas-X® Ultra Strength

Drug Facts (continued)

Directions

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*This product is not manufactured or distributed by GlaxoSmithKline Consumer Healthcare Holdings (US) LLC, owner of the registered trademark Gas-X® Ultra Strength Softgels.

ULTRA STRENGTH GAS RELIEF

simethicone capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-585
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 (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KOOR)	

Product Characteristics

Color	orange (clear)	Score	no score
Shape	CAPSULE (oval)	Size	10mm
Flavor		Imprint Code	PC3
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-585-21	50 in 1 BOTTLE; Type 0: Not a Combination Product	10/24/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M002	10/24/2017	

Labeler - Wal-Mart Stores Inc (051957769)

Establishment

Name	Address	ID/FEI	Business Operations
Humanwell PuraCap Pharmaceutical (Wuhan) Co., Ltd.		421293287	manufacture(49035-585)

Revised: 11/2024

Wal-Mart Stores Inc