GAS RELIEF- simethicone capsule, liquid filled
Advanced Rx LLC

Drug Facts

Active ingredient (in each softgel)

Simethicone 250 mg

Purpose

Antigas

Use

for the relief of pressure, bloating, and 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

- **adults and children 12 years and older:** swallow with water 1 or 2 softgels as needed after meals and at bedtime
- do not exceed 2 softgels in 24 hours except under the advice and supervision of a physician

Other information

- store at 20° to 25°C (68° to 77°F)
- protect from light, heat and moisture
- DO NOT USE IF PRINTED SEAL UNDER CAP IS MISSING OR DAMAGED

Inactive ingredients

D&C red #33, FD&C blue #1, gelatin, glycerin, purified water and white edible ink.

Questions?

call **1-800-630-8895**

Distributed by:

Advanced Rx LLC

1942 NE 163rd St North Miami Beach,

FL 33162 U.S.A.

NDC 80513-112-01

Compare to the active ingredient in Maximum Strength Gas-X ®†

HIGH POTENCY

GAS RELIEF

Simethicone 250 mg

• RELIEVES GAS FAST

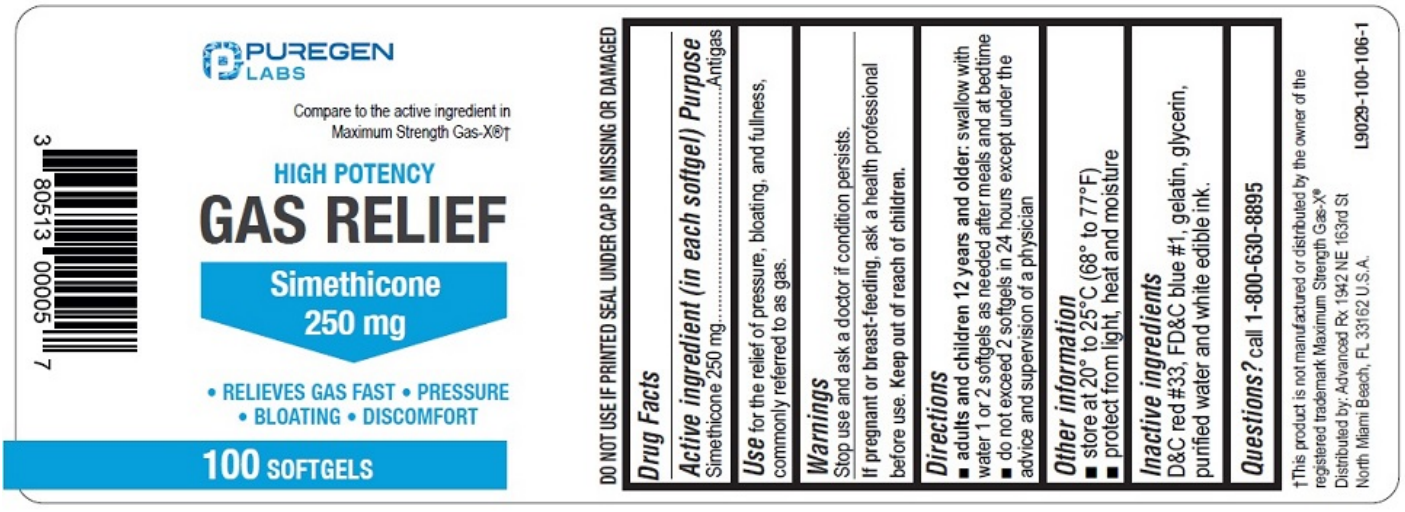
• PRESSURE

• BLOATING

• DISCOMFORT

100 SOFTGELS

†This product is not manufactured or distributed by the owner of the registered trademark Maximum Strength Gas-X ®



GAS RELIEF

simethicone capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80513-112
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	250 mg

Inactive Ingredients

Ingredient Name	Strength
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	purple (Transparent Purple)	**Score**	no score
Shape	OVAL	**Size**	13mm
Flavor		**Imprint Code**	PC31
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80513-112-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	08/30/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M002	08/30/2024	

Labeler - Advanced Rx LLC (042795108)

Revised: 8/2024 Advanced Rx LLC