GAS RELIEF ULTRA STRENGTH- simethicone capsule, liquid filled Walgreens

Drug Facts

Active ingredient (in each softgel)

Simethicone 180 mg

Purpose

Antigas

Uses

For the relief of

pressure and bloating commonly referred to as gas

Warnings

If pregnant or breast-feeding,

ask a health professional before use.

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 except under the advice and supervision of a physician

Other information

- store between 15-30°C (59-86°F)
- protect from heat and moisture

Inactive ingredients

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

Questions or comments?

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal display panel

Compare to the active ingredient in Gas-X® Ultra Strengt Softgels††

Gas Relief

SIMETHICONE 180 mg ANTIGAS

Ultra Strength

SOFTGELS

• For fast relief of pressure, bloating & fullness

††This product is not manufactured or distributed by GSK Consumer Healthcare, distributor of Gas-X® Ultra Strength Softgels.

TAMPER EVIDENT: DO NOT USE IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

DISTRIBUTED BY:WALGREEN CO.

200 WILMOT RD., DEERFIELD, IL 60015

Package Label



WALGREENS Ultra Strength Gas Relief

GAS RELIEF ULTRA STRENGTH

simethicone capsule, liquid filled

ı	Product Information			
	Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-8740
	Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
DIMETHICONE (UNII: 92RU3N3Y10) (DIMETHICONE - UNII:92RU3N3Y10)	DIMETHICONE	180 mg

Inactive Ingredients

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

Product Characteristics			
Color	orange	Score	no score
Shape	OVAL	Size	12mm
Flavor		Imprint Code	05A
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:0363-8740- 50	50 in 1 CARTON	05/29/2020		
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product			
2	NDC:0363-8740- 20	20 in 1 CARTON	05/29/2020		
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M002	05/29/2020	

Labeler - Walgreens (008965063)

Revised: 6/2024 Walgreens