SIMETHICONE- simethicone capsule, liquid filled WALMART INC.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

MAXIMUM STRENGTH GAS RELIEF

Active ingredient

Simethicone 250 mg

Purpose

Anti-gas

Uses

relieves bloating, pressure or fullness commonly referred to as gas

Warnings

Stop use and ask a doctor if

condition persists

Keep Out of Reach of Children

Directions

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

Other Information

■ store at room temperature 59°-86°F (15°-30°C)

Inactive Ingredients

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

Questions or comments?

Call toll free: 1-855-215-8180

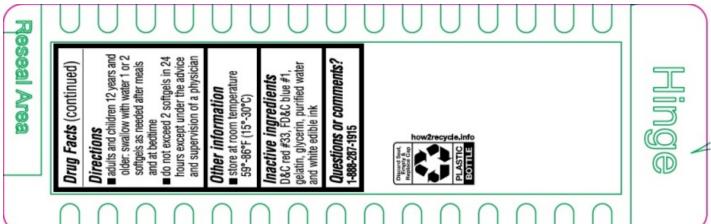
Package/Label Principal Display Panel - Principal Display Panel

Maximum Strength Gas Relief

Simethicone 250 mg 60ct

NDC: 79903-020-22





SIMETHICONE

simethicone capsule, liquid filled

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Duaduat	Information
Product	Intormation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:79903-020

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength
DIMETUICONE (LINII, ODDIDNOVIO) (DIMETUICONE	LINIII.O2DLI2NI2V1O)	DIMETHICONE	250 mg

DIMETHICONE (UNII: 92RU3N3Y10) (DIMETHICONE - UNII:92RU3N3Y10) DIMETHICONE 250 mg

Inactive Ingredients

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

Product Characteristics			
Color	PURPLE (Clear)	Score	no score
Shape	CAPSULE (Oval)	Size	13mm
Flavor		Imprint Code	PC31
Contains			

ı	Packaging			
4	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-020- 22	60 in 1 BOTTLE; Type 0: Not a Combination Product	12/18/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part332	12/18/2020	

Labeler - WALMART INC. (051957769)

Revised: 2/2021 WALMART INC.