PREMIER VALUE INFANTS GAS RELIEF- simethicone emulsion
Chain Drug Consortium

Premier Value Infants' Drops Gas Relief Drug Facts

Active ingredient (in each 0.3 mL)

Simethicone 20 mg

Purpose

Antigas

Uses

relieves the discomfort of infant gas frequently caused by air swallowing or certain formulas or foods

Warnings

Keep out of reach of children. In case of overdose get medical help or contact a poison control center right away.

Directions

- shake well before using
- all dosages may be repeated as needed, after meals and at bedtime or as directed by a physician. Do not exceed 12 doses per day.
- fill enclosed dropper to recommended dosage level and dispense liquid slowly into baby’s mouth, toward the inner cheek
- dosage can also be mixed with 1 oz. cool water, infant formula or other suitable liquids
- clean dropper after each use - replace bottle with original cap

age (yr)	weight (lb)	dose
infants under 2	under 24	0.3 mL
children over 2	over 24	0.6 mL

Other information

- **tamper evident: do not use if printed seal under cap is torn or missing**
- store at room temperature
- do not freeze
- see bottom panel for lot and expiration date

Inactive ingredients

carboxymethylcellulose sodium, citric acid, flavors, microcrystalline cellulose, polysorbate 60, potassium sorbate, purified water, sodium benzoate, sorbitan monostearate, sorbitol, xanthan gum

Principal Display Panel

COMPARE TO ACTIVE INGREDIENT IN INFANTS' MYLICON® DROPS*

Premier Value®

Infants'

Gas Relief

Non-Staining formula

Simethicone 20 mg/Antigas

- Dye Free
- No Saccharin
- No Artificial Colors
- No Artificial flavors

Syringe Enclosed. This Bottle contains 100 doses (0.3 mL/dose)

1 Fl. OZ (30ml)

100 Doses

Distributed by

Pharmacy Value Alliance, LLC

407 East Lancaster Avenue,

Wayne, PA 19087

*This product is not manufactured or distributed by Infirst Healthcare Inc., the distributor of Infants’ Mylicon® Drops.

Premier Value Infants' Gas Relief

PREMIER VALUE INFANTS GAS RELIEF

simethicone emulsion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68016-670
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIMETHICONE, UNSPECIFIED (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE, UNSPECIFIED	20 mg in 0.3 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITAN MONOSTEARATE (UNII: NVZ4I0H58X)	
SORBITOL (UNII: 506T60A25R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	WHITE (white to off white, opaque)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-670-00	1 in 1 CARTON	01/09/2015	
1		30 mL in 1 BOTTLE, DROPPER; 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	01/09/2015	

Labeler - Chain Drug Consortium (101668460)

Revised: 10/2024 Chain Drug Consortium