

**QUALITY CHOICE INFANTS DYE FREE GAS RELIEF- simethicone emulsion**  
**CHAIN DRUG MARKETING ASSOCIATION**

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**Quality Choice Infants Dye Free Gas Relief Drops**

**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.**

**Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away at 1-800-222-1222. Quick medical attention is critical even if you do not notice any signs or symptoms.**

**Directions**

- **shake well before using**
- find right dose on chart below. If possible, use weight to dose; otherwise, use age.
- only use the enclosed syringe. **Do not use other syringe, dropper, spoon or dosing device when giving this medicine to your child.**
- remove cap and insert syringe into the bottle
- pull syringe up until filled to the prescribed level. If you pass the prescribed level, simply push syringe back until you have reached the desired level. Slowly dispense the liquid into your child's mouth (toward inner cheek).
- 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.
- dosage can also be mixed with 1 oz. cool water, infant formula or other suitable liquids
- replace cap tightly to maintain child resistance
- mL = milliliter

Age (yr)	Weight (lb)	Dose (mL)
infants under 2	under 24	0.3

children over 2	over 24	0.6
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### Other information

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

### Questions or comments ?

**Call 1-866-467-2748**

### Principal Display Panel

NDC# **63868-696-30**

QUALITY CHOICE<sup>®</sup>

\*Compare to the Active Ingredient in Infants' Mylicon<sup>®</sup>

Infants' Dye-Free Gas Relief Drops

Relieves Gas Symptoms

Simethicone Antigas 20 mg

No Saccharin

No Artificial Colors

No Artificial Flavor

Non-Staining

1 Fl. OZ (30ml)

100 Doses

100% QC SATISFACTION GUARANTEED

Distributed by: C.D.M.A. Inc. ©

43157 W 9 Mile Rd

Novi, MI 48375

[www.qualitychoice.com](http://www.qualitychoice.com)

Questions: 248-449-9300

\*This product is not manufactured or distributed by Infirst Healthcare Inc., the

distributor of Infants' MYLICON<sup>®</sup> Drops.



## QUALITY CHOICE INFANTS DYE FREE GAS RELIEF

simethicone emulsion

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63868-696
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DIMETHICONE, UNSPECIFIED</b> (UNII: 92RU3N3Y1O) (DIMETHICONE, UNSPECIFIED - UNII:92RU3N3Y1O)	DIMETHICONE, UNSPECIFIED	20 mg in 0.3 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED</b> (UNII: K679OBS311)	
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POLYSORBATE 60</b> (UNII: CAL22UVI4M)	
<b>POTASSIUM SORBATE</b> (UNII: 1VPU26JZ Z 4)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SORBITAN MONOSTEARATE</b> (UNII: NVZ 4I0H58X)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	

## Product Characteristics

<b>Color</b>	white (white to off white, opaque)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-696-30	1 in 1 CARTON	12/17/2018	
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	12/17/2018	

**Labeler -** CHAIN DRUG MARKETING ASSOCIATION (011920774)

Revised: 12/2024

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