

VARIBAR THIN HONEY- barium sulfate suspension
E-Z-EM Canada Inc

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use VARIBAR THIN HONEY safely and effectively. See full prescribing information for VARIBAR THIN HONEY.

VARIBAR THIN HONEY (barium sulfate) oral suspension

Initial U.S. Approval: 2016

----- **INDICATIONS AND USAGE** -----

VARIBAR THIN HONEY is a radiopaque contrast agent indicated for use in modified barium swallow examinations to evaluate the oral and pharyngeal function and morphology in adult and pediatric patients (1)

----- **DOSAGE AND ADMINISTRATION** -----

- For oral use only – administer by syringe, spoon, or cup. The recommended dose is:
 - Adults: 5 mL
 - Pediatric patients: 1 to 3 mL
- During a single modified barium swallow examination, multiple doses may be administered
- Maximum cumulative dose: 30 mL (2)

----- **DOSAGE FORMS AND STRENGTHS** -----

Oral suspension: barium sulfate (40% w/v) supplied in a multiple-dose bottle for oral administration (3)

----- **CONTRAINDICATIONS** -----

- Known or suspected perforation of the gastrointestinal (GI) tract (4)
- Known obstruction of the GI tract (4)
- Conditions associated with high risk of GI perforation or aspiration (4)
- Known hypersensitivity to barium sulfate or any of the excipients of VARIBAR THIN HONEY (4)

----- **WARNINGS AND PRECAUTIONS** -----

- Hypersensitivity reactions: Emergency equipment and trained personnel should be immediately available (5.1)
- Intra-abdominal leakage: May occur in conditions such as GI fistula, ulcer, inflammatory bowel disease, appendicitis or diverticulitis, severe stenosis or obstructing lesions of the GI tract (5.2)
- Delayed GI transit and obstruction: Patients should maintain adequate hydration in days following barium sulfate procedure to avoid obstruction or impaction (5.3)
- Aspiration pneumonitis: Aspiration may occur during the modified barium swallow examination, monitor the patient for aspiration (5.4)

----- **ADVERSE REACTIONS** -----

Common adverse reactions include nausea, vomiting, diarrhea and abdominal cramp (6)

To report SUSPECTED ADVERSE REACTIONS, contact Bracco Diagnostics Inc at 1-800-257-5181 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

See 17 for PATIENT COUNSELING INFORMATION
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Revised: 10/2023

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* Sections or subsections omitted from the full prescribing information are not listed.

FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

VARIBAR THIN HONEY is indicated for modified barium swallow examinations to evaluate the oral and pharyngeal function and morphology in adult and pediatric patients.

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Dosing

- The recommended dose of VARIBAR THIN HONEY administered orally by syringe, spoon, or cup is:
 - Adults: 5 mL
 - Pediatric patients: 1 to 3 mL
- During a single modified barium swallow examination, multiple doses of VARIBAR THIN HONEY may be administered, to assess the patient during multiple swallows and different radiographic views.
- The maximum cumulative dose is 30 mL.

- Once opened, write the discard after date on the immediate container label. Discard any unused product after 21 days.

2.2 Important Administration Instructions

- For oral use only
- Advise patients to hydrate following the barium sulfate procedure.

3 DOSAGE FORMS AND STRENGTHS

Oral suspension: barium sulfate (40% w/v) supplied in a multiple-dose plastic bottle as a ready-to-use suspension for oral administration. Each bottle contains 250 mL of suspension.

4 CONTRAINDICATIONS

VARIBAR THIN HONEY is contraindicated in patients with:

- known or suspected perforation of the gastrointestinal (GI) tract
- known obstruction of the GI tract
- high risk of GI perforation such as those with a recent GI perforation, acute GI hemorrhage or ischemia, toxic megacolon, severe ileus, post GI surgery or biopsy, acute GI injury or burn, or recent radiotherapy to the pelvis
- high risk of aspiration such as those with known or suspected tracheo-esophageal fistula or obtundation
- known severe hypersensitivity to barium sulfate or any of the excipients of VARIBAR THIN HONEY

5 WARNINGS AND PRECAUTIONS

5.1 Hypersensitivity Reactions

Barium sulfate preparations contain a number of excipients, including natural and artificial flavors and may induce serious hypersensitivity reactions. The manifestations include hypotension, bronchospasm and other respiratory impairments, and dermal reactions including rashes, urticaria and itching. A history of bronchial asthma, atopy, food allergies, or a previous reaction to a contrast agent may increase the risk for hypersensitivity reactions. Emergency equipment and trained personnel should be immediately available for treatment of a hypersensitivity reaction.

5.2 Intra-abdominal Barium Leakage

The use of VARIBAR THIN HONEY is contraindicated in patients at high risk of perforation of the GI tract [*see Contraindications (4)*]. Administration of VARIBAR THIN HONEY may result in leakage of barium from the GI tract in the presence of conditions such as carcinomas, GI fistula, inflammatory bowel disease, gastric or duodenal ulcer, appendicitis, or diverticulitis, and in patients with a severe stenosis at any level of the GI tract, especially if it is distal to the stomach. The barium leakage has been associated with peritonitis and granuloma formation.

5.3 Delayed Gastrointestinal Transit and Obstruction

Orally administered barium sulfate may accumulate proximal to a constricting lesion of the colon, causing obstruction or impaction with development of baroliths (inspissated barium associated with feces) and may lead to abdominal pain, appendicitis, bowel obstruction, or rarely perforation. Patients with the following conditions are at higher risk for developing obstruction or baroliths: severe stenosis at any level of the GI tract, impaired GI motility, electrolyte imbalance, dehydration, on a low residue diet, taking medications that delay GI motility, constipation, pediatric patients with cystic fibrosis or Hirschsprung disease, and the elderly [see *Use in Specific Populations (8.4, 8.5)*]. To reduce the risk of delayed GI transit and obstruction, patients should maintain adequate hydration after the barium sulfate procedure.

5.4 Aspiration Pneumonitis

The use of VARIBAR THIN HONEY is contraindicated in patients with trachea-esophageal fistula [see *Contraindications (4)*]. Oral administration of barium is associated with aspiration pneumonitis, especially in patients with a history of food aspiration or with compromised swallowing mechanism. Vomiting following oral administration of barium sulfate may lead to aspiration pneumonitis.

In patients at risk for aspiration, begin the procedure with a small ingested volume of VARIBAR THIN HONEY. Monitor the patient closely for aspiration, discontinue administration of VARIBAR THIN HONEY if aspiration is suspected, and monitor for development of aspiration pneumonitis.

5.5 Systemic Embolization

Barium sulfate products may occasionally intravasate into the venous drainage of the GI tract and enter the circulation as a "barium embolus" leading to potentially fatal complications which include systemic and pulmonary embolism, disseminated intravascular coagulation, septicemia and prolonged severe hypotension. Although this complication is exceedingly uncommon after oral administration of a barium sulfate suspension, monitor patients for potential intravasation when administering barium sulfate.

6 ADVERSE REACTIONS

The following adverse reactions have been identified from spontaneous reporting or clinical studies of barium sulfate administered orally. Because the reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or to establish a causal relationship to drug exposure:

- Nausea, vomiting, diarrhea and abdominal cramping
- Serious adverse reactions and fatalities include aspiration pneumonitis, barium sulfate impaction, intestinal perforation with consequent peritonitis and granuloma formation, vasovagal and syncopal episodes

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

VARIBAR THIN HONEY is not absorbed systemically following oral administration, and maternal use is not expected to result in fetal exposure to the drug.

8.2 Lactation

Risk Summary

VARIBAR THIN HONEY is not absorbed systemically by the mother following oral administration, and breastfeeding is not expected to result in exposure of the infant to the drug.

8.4 Pediatric Use

The efficacy of VARIBAR THIN HONEY in pediatric patients is based on successful opacification of the pharynx during modified barium swallow examinations [see *Clinical Pharmacology (12.1)*]. Safety and dosing recommendations in pediatric patients are based on clinical experience.

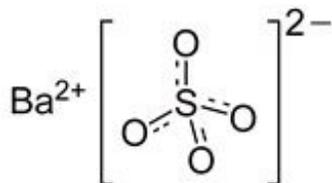
VARIBAR THIN HONEY is contraindicated in pediatric patients with trachea-esophageal fistula. [see *Contraindications (4)*]. Pediatric patients with a history of asthma or food allergies may be at increased risk for development of hypersensitivity reactions [see *Warnings and Precautions (5.1)*]. Monitor patients with cystic fibrosis or Hirschsprung disease for bowel obstruction after use [see *Warnings and Precautions (5.3)*].

8.5 Geriatric Use

Clinical studies of VARIBAR THIN HONEY did not include sufficient numbers of subjects aged 65 and over to determine whether they respond differently from younger subjects. Other reported clinical experience has not identified differences in responses between the elderly and younger patients. In general, dose selection for an elderly patient should be cautious, usually starting at the low end of the dosing range, reflecting the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other drug therapy.

11 DESCRIPTION

VARIBAR THIN HONEY (barium sulfate) is a radiographic contrast agent that is supplied as an off-white to lightly colored suspension (40% w/v) with an apple aroma for oral administration. The active ingredient barium sulfate is designated chemically as BaSO₄ with a molecular weight of 233.4 g/mol, a density of 4.5 g/cm³, and the following chemical structure:



VARIBAR THIN HONEY has a viscosity of 1500 cPs and contains the following excipients:

carboxymethylcellulose sodium, citric acid, glycerin, natural and artificial apple flavor, polysorbate 80, potassium sorbate, purified water, saccharin sodium, simethicone emulsion, sodium benzoate, sodium citrate, starch modified (from corn), xanthan gum, and xylitol.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Due to its high atomic number, barium (the active ingredient in VARIBAR THIN HONEY) is opaque to x-rays and therefore acts as a positive contrast agent for radiographic studies.

12.2 Pharmacodynamics

Barium sulfate is biologically inert and has no known pharmacological effects.

12.3 Pharmacokinetics

Under physiological conditions, barium sulfate passes through the gastrointestinal tract in an unchanged form and is absorbed only in small, pharmacologically insignificant amounts.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

No animal studies have been performed to evaluate the carcinogenic potential of barium sulfate or potential effects on fertility.

16 HOW SUPPLIED/STORAGE AND HANDLING

16.1 How Supplied

VARIBAR THIN HONEY is supplied as a suspension in a multiple-dose polyethylene bottle containing 250 mL of barium sulfate (40 % w/v).

Provided as: 12 x 250 mL bottles (NDC 32909-121-07)

16.2 Storage and Handling

Store at USP controlled room temperature 20°C to 25°C (68°F to 77°F). Protect from freezing.

Once opened, VARIBAR THIN HONEY may be used for up to 21 days when stored at USP controlled room temperature, 20°C to 25°C (68°F to 77°F).

17 PATIENT COUNSELING INFORMATION

After administration, advise patients to:

- Maintain adequate hydration [*see Dosage and Administration (2.2) and Warnings and Precautions (5.3)*].

- Seek medical attention for worsening of constipation or slow gastrointestinal passage [see *Warnings and Precautions (5.3)*].
- Seek medical attention for any delayed onset of hypersensitivity: rash, urticaria, or respiratory difficulty [see *Warnings and Precautions (5.1)*].

Rx only

Manufactured by
 EZEM Canada Inc
 Anjou (Quebec) Canada H1J 2Z4

For
 Bracco Diagnostics Inc.
 Monroe Township, NJ 08831

VARIBAR is a registered trademark of E-Z-EM, Inc.

October 2023

Varibar Thin Honey External Label

NDC 32909-121-07 12 x 250 mL

VARIBAR® THIN HONEY



Bracco Diagnostics

(BARIUM SULFATE) ORAL SUSPENSION, 40% w/v
Multiple-dose container – For oral use only
 For use in modified barium swallow examinations to evaluate the oral and pharyngeal function and morphology in adult and pediatric patients.
 Usual dosage: See prescribing information
 Each mL contains 0.4 g barium sulfate and the following inactive ingredients: carboxymethylcellulose sodium, citric acid, glycerin, natural and artificial apple flavor, polysorbate 80, potassium sorbate, purified water, saccharin sodium, simethicone emulsion, sodium benzoate, sodium citrate, starch modified (from corn), xanthan gum, xylitol.

Store at USP controlled room temperature, 20°C to 25°C (68°F to 77°F).
 Protect from freezing.

Rx only

Manufactured by
 E-Z-EM Canada Inc, Anjou, Quebec H1J2Z4, Canada
 for Bracco Diagnostics Inc., Monroe Twp., NJ 08831


 (01)30332909121076

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Lot and expiry encoding area
1" x 2"

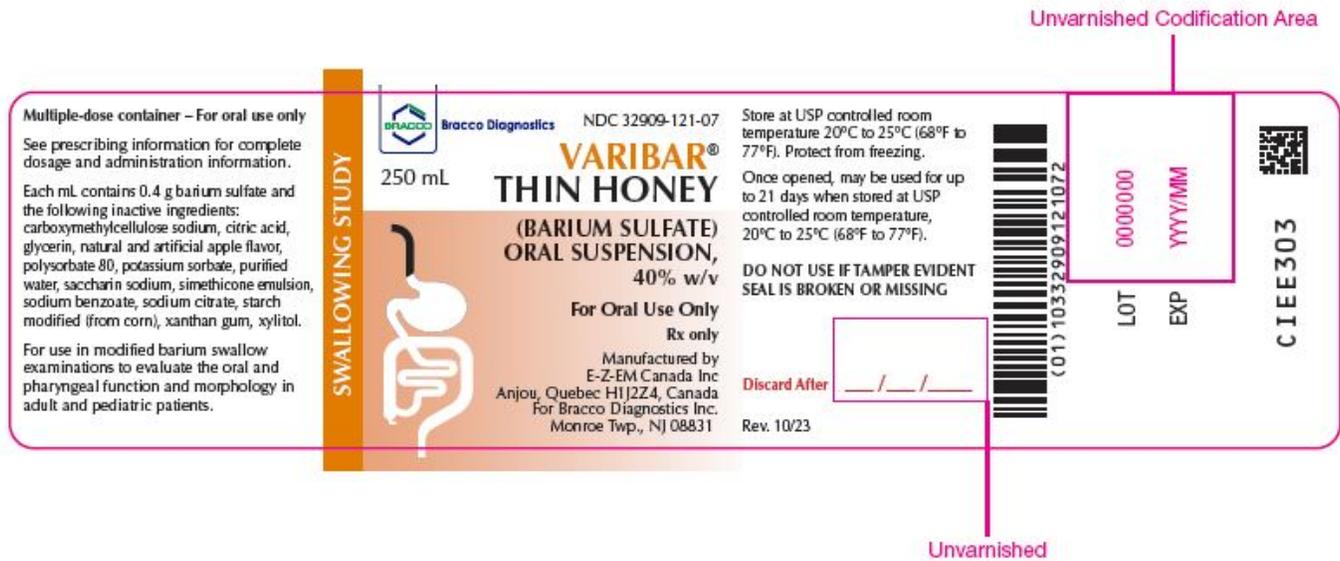
LOT

EXP.

CEEE203



Varibar Thin Honey Unit Label



VARIBAR THIN HONEY

barium sulfate suspension

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:32909-121
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Barium Sulfate (UNII: 25BB7EKE2E) (Barium Sulfate - UNII:25BB7EKE2E)	Barium Sulfate	400 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
anhydrous citric acid (UNII: XF417D3PSL)	
carboxymethylcellulose sodium (UNII: K679OBS311)	
dimethicone 350 (UNII: 2Y53S6ATLU)	
dimethicone 1000 (UNII: MCU2324216)	
glycerin (UNII: PDC6A3C00X)	
modified corn starch (1-octenyl succinic anhydride) (UNII: 461P5CJN6T)	
polysorbate 80 (UNII: 6OZP39ZG8H)	
potassium sorbate (UNII: 1VPU26JZZ4)	
silicon dioxide (UNII: ETJ7Z6XBU4)	
sodium benzoate (UNII: OJ245FE5EU)	
trisodium citrate dihydrate (UNII: B22547B95K)	
water (UNII: 059QF0KO0R)	
xanthan gum (UNII: TTV12P4NEE)	
xylitol (UNII: VCQ006KQ1E)	
saccharin sodium (UNII: SB8ZUX40TY)	

Product Characteristics

Color	WHITE	Score	
Shape		Size	
Flavor	APPLE	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:32909-121-07	12 in 1 CASE	01/03/2018	
1		250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA208143	01/03/2018	

Labeler - E-Z-EM Canada Inc (204211163)

Registrant - E-Z-EM, INC. (002041226)

Establishment

Name	Address	ID/FEI	Business Operations
E-Z-EM Canada Inc		204211163	MANUFACTURE(32909-121) , LABEL(32909-121) , ANALYSIS(32909-121) , PACK(32909-121)

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E-Z-EM Canada Inc