

**QUALITY CHOICE ULTRA STRENGTH STOMACH RELIEF- bismuth subsalicylate liquid**  
**QUALITY CHOICE (CHAIN DRUG MARKETING ASSOCIATION)**

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**Quality Choice Ultra Strength Stomach Relief 525mg**

**Active ingredient (in each 15 mL dose)**

Bismuth subsalicylate 525 mg

**Purpose**

Upset stomach reliever/Antidiarrheal

**Uses**

relieves

- travelers' diarrhea
- diarrhea
- upset stomach due to overindulgence in food and drink, including:
  - heartburn
  - indigestion
  - nausea
  - gas
  - belching
  - fullness

**Warnings**

**Reye's syndrome:** Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's syndrome, a rare but serious illness.

**Allergy alert: Contains salicylate. Do not take if you are**

- allergic to salicylates (including aspirin)
- taking other salicylate products

**Do not use**

if you have

- an ulcer
- a bleeding problem
- bloody or black stool

**Ask a doctor before use if you have**

- fever

- mucus in the stool

### **Ask a doctor or pharmacist before use if you are**

taking any drug for

- anticoagulation (thinning the blood)
- diabetes
- gout
- arthritis

### **When using this product**

a temporary, but harmless, darkening of the stool and/or tongue may occur.

### **Stop use and ask a doctor if**

- symptoms get worse or last more than 2 days
- ringing in the ears or loss of hearing occurs
- diarrhea lasts more than 2 days

### **If pregnant or breast-feeding,**

ask a health professional before use.

### **Keep out of reach of children.**

In case of overdose, get medical help or contact a Poison Control Center right away at 1-800-222-1222.

### **Directions**

- shake well before use
- Only use dose cup provided
- **adults and children 12 years and over**
  - 15 mL (1 dose) every ½ hour or 30 mL (2 doses) every hour as needed for diarrhea/traveler's diarrhea.
  - 15 mL (1 dose) every ½ hour as needed for overindulgence (upset stomach, heartburn, indigestion, nausea)
  - **do not exceed 8 doses (120 mL) in 24 hours**
  - use until diarrhea stops but not more than 2 days
- **children under 12 years:**ask a doctor
- drink plenty of clear fluids to help prevent dehydration caused by diarrhea

### **Other information**

- **each 15 mL dose cup contains:**potassium 7 mg, sodium 5 mg
- salicylate 230 mg
- low sodium
- sugar free
- protect from freezing
- avoid excessive heat (more than 104°F or 40°C)
- **TAMPER EVIDENT: Do not use if imprinted shrink band is missing or broken**

## **Inactive ingredients**

Carboxymethylcellulose sodium, D&C Red No. 22, D&C Red No. 28, flavor, gellan gum, microcrystalline cellulose, potassium hydroxide, potassium sorbate, purified water, salicylic acid, simethicone emulsion, sodium benzoate, sucralose, xanthan gum.

## **Principal Display Panel**

\*Compare to the active ingredient in **Pepto-Bismol® Ultra Strength**

QUALITY CHOICE

Ultra Strength

Stomach Relief

**Bismuth Subsalicylate 525 mg**

Upset Stomach Reliever/ Antidiarrheal

### **Relieves:**

Nausea, Heartburn, Indigestion, Upset Stomach & Diarrhea

2X CONCENTRATED FORMULA\*\*

Original Flavor

8 FL OZ (236 mL)

100% QC SATISFACTION GUARANTEED

**\*This product is not manufactured or distributed by The Procter & Gamble, distributors of Pepto-Bismol® Ultra Strength.**

**Distributed by CDMA, Inc.**

**Novi, MI 48375**

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

**Questions: 800-935-2362**



## Package Label for 12 FL OZ 354 ml



**QUALITY CHOICE ULTRA STRENGTH STOMACH RELIEF**  
bismuth subsalicylate liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:83324-337
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>BISMUTH SUBSALICYLATE</b> (UNII: 62TEY51RR1) (BISMUTH CATION - UNII:Z59CD1I8YE, SALICYLIC ACID - UNII:O414PZ4LPZ)	BISMUTH SUBSALICYLATE	525 mg in 15 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED</b> (UNII: K679OBS311)	
<b>D&amp;C RED NO. 22</b> (UNII: 1678RKX8RT)	
<b>D&amp;C RED NO. 28</b> (UNII: 7671P0Y5NH)	
<b>GELLAN GUM (HIGH ACYL)</b> (UNII: W1L7G7ROMD)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POTASSIUM HYDROXIDE</b> (UNII: WZH3C48M4T)	
<b>POTASSIUM SORBATE</b> (UNII: 1VPU26JZZ4)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SALICYLIC ACID</b> (UNII: O414PZ4LPZ)	
<b>DIMETHICONE, UNSPECIFIED</b> (UNII: 92RU3N3Y1O)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	

**Product Characteristics**

<b>Color</b>	pink (viscous)	<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:83324-337-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/07/2025	
2	NDC:83324-337-12	354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/22/2025	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M008	07/07/2025	

Revised: 9/2025

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