# STOMACH RELIEF ULTRA- bismuth subsalicylate liquid QUALITY CHOICE (Chain Drug Marketing Association)

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#### **Drug Facts**

## Active ingredient (in each 15 mL)

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
  - o gas
  - indigestion
  - nausea
  - fullness
  - belching

# Warnings

**Reye's syndrome**: Children and teenagers who have or are recovering from chicken pox or flu-like symptoms ahould 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

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

#### 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 childen.

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

#### **Directions**

- do not take more than 8 doses (120 mL) in 24 hours
- use until diarrhea stops but not more than 2 days
- drink plenty of clear fluids to help prevent dehydration cause by diarrhea
- mL = mililiter
- shake well before using
- measure only with dosing cup provided. Do not use any other dosing device
- keep dosing cup with product
  - adults and children12 years and over:
  - 15 mL (1 dose) every 1/2 hour or 30 mL (2 doses) every hour as needed for diarrhea/traveler's diarrhea
  - 15 mL (1 dose) every 1/2 hour as needed for overindulgence (upset stomach, heartburn, indigestion,nausea)
- children under 12 years of age: ask a doctor

#### Other information

each 15 mL contains: sodium 6 mg
each 15 mL contains: salicylate 227 mg

- low sodium
- keep tightly closed

- protect from freezing
- avoid excessive heat (over 104°F or 40°C)

## **Inactive ingredients**

benzoic acid, D&C red #22, D&C red #28, flavor, magnesium aluminum silicate, methylcellulose, purified water, saccharin sodium, salicylic acid, sodium salicylate, sorbic acid

#### Questions or comments?

Call **1-800-935-2362** Monday-Friday 9AM-5PM EST

#### **Principal Display Panel**

\*Compare to the Active Ingredient IN Pepto-Bismol® Ultra

Ultra

#### Stomach Relief

Stomach Relief

Bismuth subsalicylate | Upset Stomach Reliever/Antidiarrheal

For Relief of:

Heartburn

Indigestion

Nausea

**Upset Stomach** 

Diarrhea

# 2x strength per ouncet

Alcohol free

Sugar free

FL OZ (mL)

\*This product is not manufactured or distributed by The Procter & Gamble Company. Pepto-Bismol® is a registered trademark of The Procter & Gamble Company.

# TAMPER EVIDENT; DO NOT USE IF PRINTED SAFETY SEAL AROUND DOSAGE CUP OR UNDER CAP IS BROKEN OR MISSING.

Distributed by C.D.M.A., Inc.©

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www.qualitychoice.com

#### Package Label



#### **OUALITY CHOICE Ultra Stomach Relief**

**Purpose** 

Antidiarrheal

#### STOMACH RELIEF ULTRA

bismuth subsalicylate liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-814
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
<b>BISMUTH SUBSALICYLATE</b> (UNII: 62TEY51RR1) (BISMUTH CATION - UNII:ZS9CD1I8YE, SALICYLIC ACID - UNII:O414PZ4LPZ)	BISMUTH SUBSALICYLATE	525 mg in 15 mL	

Inactive Ingredients		
Ingredient Name	Strength	
BENZOIC ACID (UNII: 85KN0B0MIM)		
D&C RED NO. 22 (UNII: 1678RKX8RT)		
<b>D&amp;C RED NO. 28</b> (UNII: 767IP0Y5NH)		
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)		
WATER (UNII: 059QF0KO0R)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		

SALICYLIC ACID (UNII: O414PZ4LPZ)	
SODIUM SALICYLATE (UNII: WQ1H85SYP)	
SORBIC ACID (UNII: X045W)989B)	
METHYLCELLULOSE (1500 CPS) (UNII: PONTE48364)	

l	Packaging			
	# Item Cod	le Package Description	Marketing Start Date	Marketing End Date
	1 NDC:63868- 814-08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/26/2021	

Marketing Information			
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
M008	03/26/2021		
	Application Number or Monograph Citation	Application Number or Monograph Marketing Start Citation Date	

Labeler - QUALITY CHOICE (Chain Drug Marketing Association) (011920774)

Revised: 5/2024 QUALITY CHOICE (Chain Drug Marketing Association)