ULTRA STRENGTH STOMACH RELIEF - bismuth subsalicylate 525 mg liquid Retail Business Services, LLC.

Ultra Strength Stomach Relief

ACTIVE INGREDIENT(in each 15 mL)

Bismuth subsalicylate 525 mg

PURPOSE

Upset stomach reliever and anti-diarrheal

USE(S)

relieves:

- diarrhea
- heartburn
- indigestion
- nausea
- upset stomach associated with these symptoms

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
- bloody or black stool
- a bleeding problem

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 of 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 DOCTOR IF

- symptoms get worse
- 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 immediately.

DIRECTIONS

- shake well before use
- mL = milliliter
- TBSP = tablespoon
- adults and children 12 years and over: 1 dose (1 TBSP or 15 mL) every 1/2 hour as needed
- 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 fluids to help prevent dehydration caused by diarrhea

OTHER INFORMATION

- each 15 mL or 1 TBSP contains:
- potassium 15 mg
- salicylate 236 mg
- sodium 5 mg
- protect from freezing

- avoid excessive heat (over 104°F or 40°C)
- dosage cup provided

INACTIVE INGREDIENTS

benzoic acid, D&C red # 22, D&C red # 28, flavor, hydroxyethyl cellulose, potassium hydroxide, purified water, saccharin sodium, salicylic acid, simethicone, xanthan gum

PRINCIPAL DISPLAY PANEL

CAREOne

NDC 72476-528-04

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

Bismuth Subsalicylate 525 mg

Upset stomach reliever/Antidiarrheal

Original Flavor

SOOTHING RELIEF FOR:

Upset stomach, Indigestion, Nausea, Heartburn, Diarrhea

PROTECTIVE COATING

8 FL OZ (236 mL)





Drug Facts (continued) Directions ■ shake well before use

■ mL = milliliter ■ TBSP = tablespoon ■ adults and children 12 years and over: 1 dose (1 TBSP or 15 mL) every 1/2 hour as needed 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 or 1 TBSP contains potassium 15 mg
 salicylate 236 mg
 sodium 5 mg ■ protect from freezing. ■ avoid excessive heat (over 104°F or 40°C). ■ dosage cup provided

Inactive ingredients benzoic acid, D&C red # 22 D&C red # 28, flavor, hydroxyethyl cellulose, potassium hydroxide, purified water, saccharin sodium, salicylic acid, simethicone, xanthan gum.

This product is not manufactured or distributed by Procter & Gamble, Inc., the distributor of Pepto-Bismol® Ultra Strength

DISTRIBUTED BY: ADUSA DISTRIBUTION, LLC SALISBURY, NC 28147

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T-0190 0725 營 53041 MFR#

ULTRA STRENGTH STOMACH RELIEF

bismuth subsalicylate 525 mg liquid

Product Information

HUMAN OTC DRUG NDC:72476-528 **Product Type** Item Code (Source)

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (SALICYLIC ACID -**BIS MUTH** 525 mg UNII:O414PZ4LPZ) **SUBSALICYLATE** in 15 mL

Inactive Ingredients	
Ingredient Name	Strength
BENZOIC ACID (UNII: 8SKN0B0MIM)	
D&C RED NO. 22 (UNII: 1678RKX8RT)	
D&C RED NO. 28 (UNII: 767IP0Y5NH)	
POTASSIUM HYDROXIDE (UNII: WZ H3C48M4T)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
HYDROXYETHYL CELLULOSE (1500 MPA.S AT 1%) (UNII: L605B5892V)	
DIMETHICONE (UNII: 92RU3N3Y10)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics			
Color	PINK	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:72476-528- 04	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/21/2025	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M008	07/21/2025		

Labeler - Retail Business Services, LLC. (967989935)

Establishment				
Name	Address	ID/FEI	Business Operations	
Guardian Drug Company		119210276	MANUFACTURE(72476-528)	

Revised: 7/2025 Retail Business Services, LLC.