

**SIGNATURE CARE ULTRA STRENGTH STOMACH- bismuth
subsalicylate suspension
SAFEWAY**

Signature SELECT Ultra Strength Stomach Relief Drug Facts

Active ingredient (in each 15 mL dose)

Bismuth subsalicylate 525 mg

Purposes

Upset stomach reliever and 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 or 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 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

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

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

Questions or comments?

1-866-467-2748

Do not use if imprinted shrink band is missing or broken.

Principal Display Panel

Signature SELECT™

NDC 21130-193-12

COMPARE TO the active ingredient in Pepto-Bismol® *

STOMACH RELIEF

ULTRA STRENGTH

ORIGINAL FLAVOR

Bismuth Subsalicylate 525 MG

Upset Stomach Reliever/Antidiarrheal

Relief of:

- Nausea, Heartburn, Indigestion, Upset Stomach, Diarrhea

12 FL. OZ. (354 mL)

2X CONCENTRATED FORMULA**

Distributed by: BETTER LIVING BRANDS LLC

P.O. BOX 99

PLEASANTON, CA 94566-0009

‡1-888-723-3929

****vs Original Pepto Bismol®**

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



SIGNATURE CARE ULTRA STRENGTH STOMACH

bismuth subsalicylate suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-193
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
D&C RED NO. 22 (UNII: 1678RKX8RT)	
D&C RED NO. 28 (UNII: 7671P0Y5NH)	
GELLAN GUM (LOW ACYL) (UNII: 7593U09I4D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
POTASSIUM SORBATE (UNII: 1VPU26JZ Z4)	
WATER (UNII: 059QF0KO0R)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
DIMETHICONE, UNSPECIFIED (UNII: 92RU3N3Y1O)	

SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	PINK (viscous)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-193-12	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/02/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M008	02/02/2026	

Labeler - SAFEWAY (009137209)

Revised: 2/2026

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