

BISMUTINA- bismuth subsalicylate liquid
Menper Distributors, Inc.

Bismutina

Active Ingredients

Bismuth Subsalicylate 525 mg

Purpose

Upset Stomach Reliever and Antidiarrheal

Uses

relieves

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

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 for more than two 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 accidental overdose, get medical help or contact a Poison Control Center right away.

Directions

- drink plenty of clear fluids to help prevent dehydration cause by diarrhea
- shake well before use
- **adults and children 12 years of age and over:**
- 2 tablespoonfuls (30 mL) every 1/2 hour or 4 tablespoonfuls (60 mL) every hour as needed for diarrhea/traveler's diarrhea
- 2 tablespoonfuls (30 mL) every 1/2 hour as needed for overindulgence (upset stomach, heartburn, indigestion, nausea)
- do not exceed 8 doses (16 tablesproonfuls or 240 mL) in 24 hours
- use until diarrhea stops but not more than 2 days
- **Children under 12 years:** ask a doctor

Inactive ingredients

benzoic acid, citric acid, FD&C yellow #5, flavor, salicylic acid, sodium citrate, sodium salicylate, sorbic acid, sucralose, water, xanthan gum

Questions? call 1-800-560-5223 M-F 9 AM to 5 PM Eastern Time



BISMUTINA
(Bismuth Subsalicylate)

**Upset Stomach Reliever/Antidiarrheal
Protective Coating Action**

Relief for:

- Heartburn
- Indigestion
- Upset Stomach
- Nausea
- Diarrhea

8 FL OZ (240 mL)

Drug Facts

Active ingredient (in each 2 tablespoon (30 mL) dose) Purposes
Bismuth subsalicylate 525 mg.....Upset Stomach Reliever
and Antidiarrheal

Uses relieves ■ traveler's diarrhea ■ diarrhea
■ upset stomach due to overindulgence in food and drink, including:
■ heartburn ■ indigestion ■ nausea ■ gas ■ belching

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Drug Facts (continued)

Other information
■ each 2 tablespoonfuls (30 mL) contains: salicylate 249 mg, and sodium 22 mg
■ sugar free ■ protect from freezing ■ avoid excessive heat (more than 104° F or 40° C)
■ **Tamper-Evident:** Do not use if printed cello-band is torn, broken or missing.

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Distributed by:
MENPER DISTRIBUTORS, INC.
11421 NW 107 St.
Suite #24
Miami, FL 33178

Rev. 1



BISMUTINA

bismuth subsalicylate liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53145-475
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII:O414PZ4LPZ)	BISMUTH SUBSALICYLATE	525 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
BENZOIC ACID (UNII: 8SKN0B0MIM)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM SALICYLATE (UNII: WQ1H85SYP)	
SORBIC ACID (UNII: X045WJ989B)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
WATER (UNII: 059QF0KO0R)	

XANTHAN GUM (UNII: TTV12P4NEE)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53145-475-08	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M024	04/01/2021	

Labeler - Menper Distributors, Inc. (101947166)

Establishment

Name	Address	ID/FEI	Business Operations
RNV, LLC		118917568	manufacture(53145-475)

Revised: 7/2025

Menper Distributors, Inc.