# DIGESTIVE RELIEF- bismuth subsalicylate tablet, film coated SAFEWAY

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#### Digestive Relief Bismuth Subsalicylate 262 mg

### **Drug Facts**

## Active ingredient (in each caplet)

Bismuth subsalicylate 262 mg

#### Purpose

Upset stomach reliever and anti-diarrheal

#### Uses

relieves:

- travelers' diarrhea
- diarrhea

upset stomach due to overindulgence in food and drink, including: ■ heartburn ■ indigestion ■ nausea ■ gas ■ belching ■ fullness

**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 immediately at 1-800-222-1222

#### **Directions**

- swallow with water, do not chew
- adults and children 12 years and older: 2 caplets every 1/2 to 1 hour as needed
- do not exceed 8 doses (16 caplets) 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 caplet contains:calcium 27 mg, sodium 3 mg
- salicylate 99 mg
- low sodium
- sugar free
- store at room temperature

#### Questions or comments?

1-866-467-2748

# **Inactive Ingredients**

calcium carbonate, crospovidone, D&C Red No. 27, magnesium stearate, maltodextrin, microcrystalline cellulose, sodium lauryl sulfate.

#### NDC# 21130-125-40

Compare to the active ingredient Pepto-Bismol®\*

# Digestive Relief

Bismuth Subsalicylate 262 mg

Upset Stomach Reliever/ Antidiarrheal

5 Symptoms Digestive Relief

• Heartburn

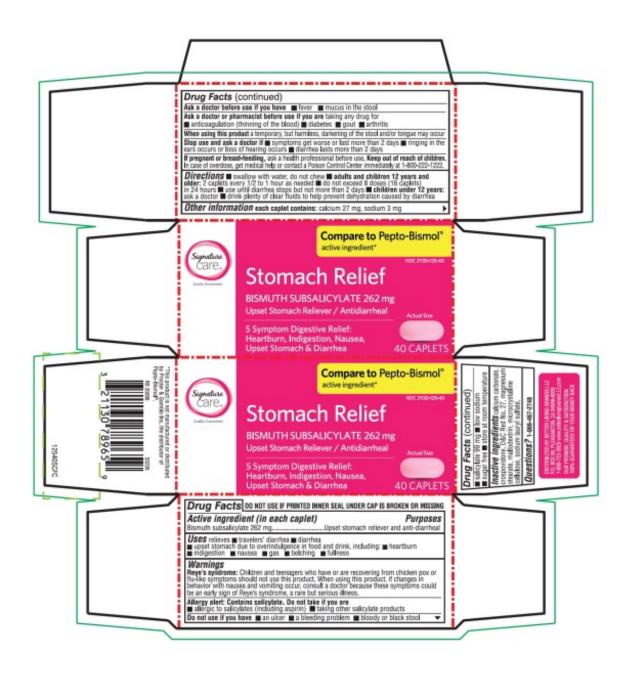
- Indigestion
- Nausea
- Upset Stomach
- Diarrhea

#### **40 CAPLETS**

# TAMPER EVIDENT: DO NOT USE IF PRINTED INNER SEAL UNDER CAP IS BROKEN OR MISSING

#### Distributed by:

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



# **DIGESTIVE RELIEF**

bismuth subsalicylate tablet, film coated

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<b>Product</b>		Ialiuii

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-125
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Route of Administration ORAL

# Active Ingredient/Active Moiety Ingredient Name BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII: 0414PZ4LPZ) BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII: 0414PZ4LPZ) BISMUTH SUBSALICYLATE

Inactive Ingredients		
Ingredient Name	Strength	
CALCIUM CARBONATE (UNII: H0G9379FGK)		
CROSPOVIDONE (120 .MU.M) (UNII: 68401960MK)		
<b>D&amp;C RED NO. 27</b> (UNII: 2LRS185U6K)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MALTODEXTRIN (UNII: 7CVR7L4A2D)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
SODIUM LAURYL SULFATE (UNII: 368GB5141J)		

Product Characteristics			
Color	pink	Score	no score
Shape	OVAL (Caplet)	Size	16mm
Flavor		Imprint Code	RP125
Contains			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:21130-125-	1 in 1 CARTON	10/05/2020	
1	40 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M008	10/05/2020	

# **Labeler -** SAFEWAY (009137209)

Revised: 10/2025 SAFEWAY