

**STOMACH RELIEF- bismuth subsalicylate tablet**  
**CHAIN DRUG MARKETING ASSOCIATION INC**

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**Quality Choice 44-346 Delisted**

***Active ingredient (in each caplet)***

Bismuth subsalicylate 262 mg

***Purpose***

Upset stomach reliever/antidiarrheal

***Uses***

relieves:

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

***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

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

## **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
- 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.

## ***Directions***

- **do not take more than directed**
- swallow with water; do not chew
- drink plenty of clear fluids to help prevent dehydration caused by diarrhea
- adults and children 12 years and over: 2 caplets every 1/2 to 1 hour as needed. Do not exceed 8 doses (16 caplets) in 24 hours.
- do not use for more than 2 days unless directed by a doctor
- use until diarrhea stops, but not more than 2 days
- children under 12 years: ask a doctor

## ***Other information***

- **each caplet contains:** calcium 20 mg, salicylate 103 mg
- store at 25°C (77°F); excursions permitted between 15°C-30°C (59°F-86°F)
- avoid excessive heat
- see end flap for expiration date and lot number

## ***Inactive ingredients***

calcium carbonate, corn starch, D&C red #27 aluminum lake, D&C red #30 aluminum lake, magnesium stearate, mannitol, microcrystalline cellulose, povidone, silicon dioxide, sodium starch glycolate, stearic acid

**Questions or comments?**

**1-800-426-9391**

***Principal display panel***

**QC®**  
QUALITY  
CHOICE

NDC 63868-691-40

\*Compare to the  
Active Ingredient in  
Pepto-Bismol®

**Stomach Relief**  
**Bismuth Subsalicylate 262 mg**  
**Upset Stomach Reliever/Antidiarrheal**

Relieves Nausea, Heartburn, Indigestion,  
Upset Stomach, Diarrhea

actual size

**40** Caplets

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY  
SEAL UNDER CAP IS BROKEN OR MISSING**

\*This product is not manufactured or distributed by The  
Procter & Gamble Company, owner of the registered  
trademark Pepto-Bismol®.  
50844 REV0521A34610

SATISFACTION  
GUARANTEED  
100% **QC**

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Novi, MI 48375  
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Questions: 800-935-2362



**Quality Choice 44-346**

**STOMACH RELIEF**

bismuth subsalicylate tablet

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63868-691
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>BISMUTH SUBSALICYLATE</b> (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII:O414PZ4LPZ, BISMUTH CATION - UNII:ZS9CD1I8YE)	BISMUTH SUBSALICYLATE	262 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>CALCIUM CARBONATE</b> (UNII: H0G9379FGK)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>D&amp;C RED NO. 27 ALUMINUM LAKE</b> (UNII: ZK64F7XSTX)	
<b>D&amp;C RED NO. 30 ALUMINUM LAKE</b> (UNII: GE75M6YV5W)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MANNITOL</b> (UNII: 3OWL53L36A)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>SODIUM STARCH GLYCOLATE TYPE A POTATO</b> (UNII: 5856J3G2A2)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	

### Product Characteristics

<b>Color</b>	pink	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	16mm
<b>Flavor</b>		<b>Imprint Code</b>	44;346
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-691-40	1 in 1 CARTON	03/31/2021	08/31/2026
1		40 in 1 BOTTLE, PLASTIC; 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	03/31/2021	08/31/2026

**Labeler** - CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(63868-691)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(63868-691)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(63868-691)

Revised: 10/2025

CHAIN DRUG MARKETING ASSOCIATION INC