

PEPTO BISMOL DIARRHEA LIQUICAPS- bismuth subsalicylate capsule, liquid filled
The Procter & Gamble Manufacturing Company

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Pepto Bismol®

Diarrhea

Drug Facts

Active ingredient (in each LiquiCap)

Bismuth subsalicylate 262 mg

Purpose

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 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 right away.

Directions

- swallow with water, do not chew
- adults and children 12 years and over:
 - 2 LiquiCaps every ½ hour or 4 LiquiCaps every hour as needed for diarrhea
 - 2 LiquiCaps every ½ hour to 1 hour as needed for overindulgence (upset stomach, heartburn, indigestion, nausea)
- do not exceed 8 doses (16 LiquiCaps) 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 LiquiCap contains:
 - salicylate 199 mg
 - low sodium
- do not exceed 25°C

Inactive ingredients

D&C red No. 33, gelatin, glycerin, polyethylene glycol, pharmaceutical ink, polysorbate 20, povidone, sorbitol sorbitan solution, titanium dioxide

Questions?

1-800-717-3786

MADE IN CANADA

DIST. BY PROCTER & GAMBLE,
CINCINNATI OH 45202

PRINCIPAL DISPLAY PANEL - 48 LiquiCap Carton

P epto®

Diarrhea

Bismuth Subsalicylate

Antidiarrheal

LIQUICAPS

RELIEVES

DIARRHEA

48 LiquiCaps™

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Warnings

Directions

Other Important Information

Questions? Ask a Doctor

Drug Facts (continued)

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Drug Facts (continued)

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Pepto Bismol Diarrhea LIQUICAPS

48 LiquiCaps™

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Pepto Bismol Diarrhea LIQUICAPS

Pepto Bismuth Subsalicylate Antidiarrheal
Diarrhea LIQUICAPS



RELIEVES DIARRHEA

48 LiquiCaps™

Pepto Bismol Diarrhea LIQUICAPS

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Also sold as Pepto Bismol® LiquiCaps
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PEPTO BISMOL DIARRHEA LIQUICAPS

bismuth subsalicylate capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37000-537
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII:O414PZ4LPZ)	BISMUTH SUBSALICYLATE	262 mg

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SORBITOL (UNII: 506T60A25R)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
GELATIN (UNII: 2G86QN327L)	
POVIDONE (UNII: FZ989GH94E)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	

Product Characteristics

Color	pink	Score	no score
Shape	CAPSULE	Size	14mm
Flavor		Imprint Code	PB
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37000-537-12	1 in 1 CARTON	03/01/2019	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:37000-537-24	2 in 1 CARTON	03/01/2019	
2		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:37000-537-48	4 in 1 CARTON	03/01/2019	
3		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part335	03/01/2018	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

