STOMACH RELIEF- bismuth subsalicylate suspension Consumer Value Products, Inc.

STOMACH RELIEF

Drug Facts

Active ingredient (in each 30 mL dose cup or 2 tablespoons)

Bismuth subsalicylate 525 mg

Purposes

Bismuth subsalicylate Upset stomach reliever and antidiarrheal

Uses relieves ■ travelers' diarrhea ■ diarrhea ■ upset stomach due to overindulgence of food and drink including:

■ heart burn ■ 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 ■ black or bloody stool

Ask a doctor before use if you have

■ fever ■ mucus in stool

Ask a doctor or pharmacist before use if you are

taking any drug for ■ anticoagulation (thinning of the blood) ■ diabetes ■ gout ■ arthritis

When using this product a temporary, but harmless, darkening of the stool and/or tongue may occur

Slop use and ask a doctor if

- diarrhea lasts more than 2 days
- symptoms get worse or last more than 2 days
- ringing in the ears or loss of hearing occurs

If pregnant or breast feeding, ask 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

- Shake well before use
- use dose cup or tablespoon (TBSP)
- adults and children 12 years and over: 1 dose (30 mL or 2 TBSP) every 1/2 to 1 hour as needed
- do not exceed 8 doses (240 mL or 16 TBSP) 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 30 mL dose

contains: sodium 13 mg, salicylate 256 mg

- protect from freezing
- avoid excessive heat (over 104°F or 40°C)
- Low sodium

Inactive ingredients benzoic acid, D&C red #22, D&C red # 28, flavor, purified water, saccharin sodium, salicylic acid, sodium salicylate, xantham gum

Questions? 1-877-798-5944

Principal Display Panel

CVP®

HEALTH

*COMPARE TO PEPTO-BISMOL® ACTIVE INGREDIENT

Stomach Relief

PROTECTIVE COATING ACTION Bismuth Subsalicylate

SOOTHING RELIEF FOR:

UPSET STOMACH
HEARTBURN
INDIGESTON
NAUSEA
COMMON DIARRHEA

8 FL OZ (237 mL)

TAMPER EVIDENT: Do not use if imprinted shrinkband is missing or broken

^{*}This product is not manufactured or distributed by Procter & Gamble,

Inc., the distributor of Pepto-Bismol™.

DISTRIBUTED BY CONSUMER VALUE PRODUCTS, INC. P.O. BOX 6115 TEMPLE, TX 76502

Made in USA LR-102 REV01

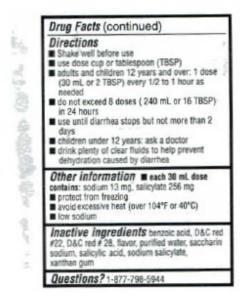
CVPproducts.com

7 61706 16202 6

PEEL

HERE





Stomach Relief by Consumer Value Products, Inc.

STOMACH RELIEF

bismuth subsalicylate suspension

Droduct	Inform	ation
Product		ativii

Product Type HUMAN OTC DRUG Item Code (Source) NDC:57243-557

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII: 0414PZ 4LPZ) BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (SALICYLIC ACID - SUBSALICYLATE in 30 mL

Inactive Ingredients		
Ingredient Name	Strength	
BENZOIC ACID (UNII: 85KN0B0MIM)		
D&C RED NO. 22 (UNII: 1678RKX8RT)		
D&C RED NO. 28 (UNII: 767IP0Y5NH)		
WATER (UNII: 059QF0KO0R)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		
SALICYLIC ACID (UNII: O414PZ4LPZ)		
SODIUM SALICYLATE (UNII: WQ1H85SYP)		
XANTHAN GUM (UNII: TTV12P4NEE)		

Product Characteristics				
Color	pink	Score		
Shape		Size		
Flavor	MINT	Imprint Code		
Contains				

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:57243-557- 26	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2018	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M008	10/01/2018		

Registrant - AptaPharma Inc. (790523323)

Establishment				
Name	Address	ID/FEI	Business Operations	
AptaPharma Inc.		790523323	manufacture(57243-557)	

Revised: 12/2023 Consumer Value Products, Inc.