

**PREMIER VALUE DIARRHEA RELIEF VANILLA FLAVOR- bismuth
subsalsicylate suspension
PHARMACY VALUE ALLIANCE, LLC**

Premier Value Diarrheal Relief

Drug Facts

Active Ingredient (per 15 mL)

Bismuth Subsalsicylate 262 mg

Purposes

Anti-diarrheal/Upset stomach reliever

Uses

relieves

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

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

- a sodium-restricted diet
- fever
- mucus in the stool

Ask a doctor or pharmacist before use if you are taking any drug for

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

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 at 1-800-222-1222.

Directions

- **shake well immediately before each use**
- **adults and children 12 years of age and older:** 30 ml or 2 tablespoonful
- for accurate dosing, use convenient pre-measured dose cup
- repeat dose every 1/2 hour to 1 hour as needed
- do not exceed 8 doses 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 15mL tablespoon contains: **sodium 10 mg**
- each 15mL tablespoon contains: **salicylate 130 mg**
- **do not use if printed inner seal is broken or missing**
- **store at room temperature**

Inactive ingredients

caramel, carboxymethylcellulose sodium, microcrystalline cellulose, natural and artificial flavor, potassium sorbate, salicylic acid, simethicone emulsion, sucralose, sucrose, water, xanthan gum

Questions or comments?

1-866-467-2748

Principal Display Panel

Premier Value®

NDC 68016-317-12

COMPARE TO THE ACTIVE INGREDIENT IN KAOPECTATE®

Diarrhea Relief

VANILLA

Regular flavor

Bismuth Subsalicylate, 262 mg

- Anti-Diarrheal
- Upset Stomach Reliever

Effective Diarrhea Relief

Restores Natural Balance

Contains Salicylates

12 FL OZ (355 mL)

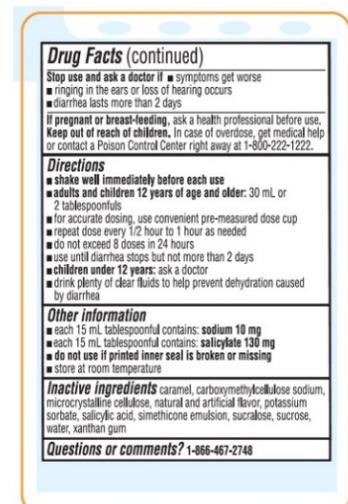
Distributed by:

Pharmacy Value Alliance, LLC

407 East Lancaster Avenue,

Wayne, PA 19087

*This product is not manufactured or distributed by Chattem Inc., the distributor of Kaopectate®.



PREMIER VALUE DIARRHEA RELIEF VANILLA FLAVOR

bismuth subsalicylate suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-317
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Bismuth subsalicylate (UNII: 62TEY51RR1) (Salicylic acid - UNII:O414PZ4LPZ, Bismuth cation - UNII:ZS9CD1I8YE)	Bismuth subsalicylate	262 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
CARAMEL (UNII: T9D99G2B1R)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
salicylic acid (UNII: O414PZ4LPZ)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SUCROSE (UNII: C151H8M554)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	WHITE	Score	
Shape		Size	
Flavor	VANILLA	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-317-12	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/23/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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OTC Monograph Drug M008

04/23/2019

Labeler - PHARMACY VALUE ALLIANCE, LLC (101668460)

Revised: 11/2023

PHARMACY VALUE ALLIANCE, LLC