KAOPECTATE VANILLA FLAVOR ANTI DIARRHEAL- bismuth subsalicylate liquid Kramer Laboratories

Drug Facts

Active ingredient (per 15 mL)	Purposes
Bismuth subsalicylate 262 mg	Anti- diarrheal Upset stomach reliever

Uses relieves:

- traveler's 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

• 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.

Directions

- shake well immediately before each use
- only use pre-measured dose cup
- adults and children 12 years of age and older:
 - 1 dose (30 mL) every 1/2 hour to 1 hour as needed.
 - do not exceed 8 doses (240 mL) 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 15 mL dose cup contains: sodium 6 mg and salicylate 135mg
- store at room temperature 20°-25° C (68°-77° F)

Inactive Ingredients

carboxymethylcellulose sodium, flavor, microcrystalline cellulose, purified water, sodium salicylate, sorbic acid, sucrose, xanthan gum

Principal Display Panel NEW IMPROVED TASTE!

Kaopectate[®] Bismuth Subsalicylate 262 mg

• Anti-Diarrheal • Upset Stomach Reliver

Diarrhea & **Upset Stomach**

- ✓ Begins controlling symptoms from the first dose
 ✓ Quickly relieves urgency, gas, and cramping
 ✓ Effective on diarrhea from bacteria,
- viruses, and other causes

11 fl oz (325mL)

Vanilla Flavor

680VAN11KA0LF



Kaopectate® Vanilla 11oz

Do not use if inner seal is broken or missing

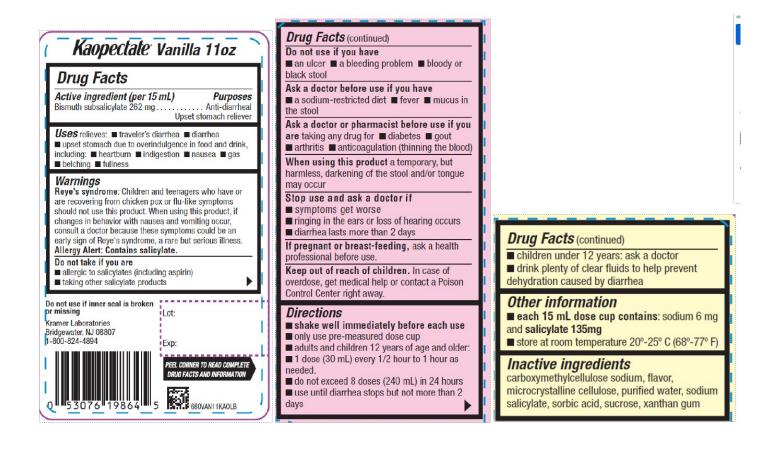
Kramer Laboratories Bridgewater, NJ 08807 1-800-824-4894

Lot:

Exp:

PEEL CORNER TO READ COMPLETE DRUG FACTS AND INFORMATION

680VAN11KA0LB



KAOPECTATE VANILLA FLAVOR ANTI DIARRHEAL

bismuth subsalicylate liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55505-198
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Bismuth Subsalicylate (UNII: 62TEY51RR1) (Bismuth Cation - UNII:ZS9CD18YE)	Bis muth Subsalicylate	262 mg in 15 mL	

Inactive Ingredients				
Ingredient Name	Strength			
Carboxymethylcellulose Sodium, Unspecified (UNII: K6790BS311)				
Microcrystalline Cellulose (UNII: OP1R32D61U)				
Sodium Salicylate (UNII: WQ1H85SYP)				
Sorbic Acid (UNII: X045WJ989B)				
Sucrose (UNII: C151H8M554)				
Water (UNII: 059QF0KO0R)				
Xanthan Gum (UNII: TTV12P4NEE)				

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	VANILLA	Imprint Code	
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55505-198- 36	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2020	
2	NDC:55505-198- 64	325 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M008	10/01/2020	

Labeler - Kramer Laboratories (122720675)

Revised: 11/2023 Kramer Laboratories