KAOPECTATE PEPPERMINT 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 20º-25º C (68º-77º F)

Inactive Ingredients

carboxymethylcellulose sodium, FD&C Red No. 40, 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 Reliever

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)

Peppermint Flavor

681PPT11KA0 F

Kaopectate Peppermint 11 oz

Do not use if inner seal is broken or missing

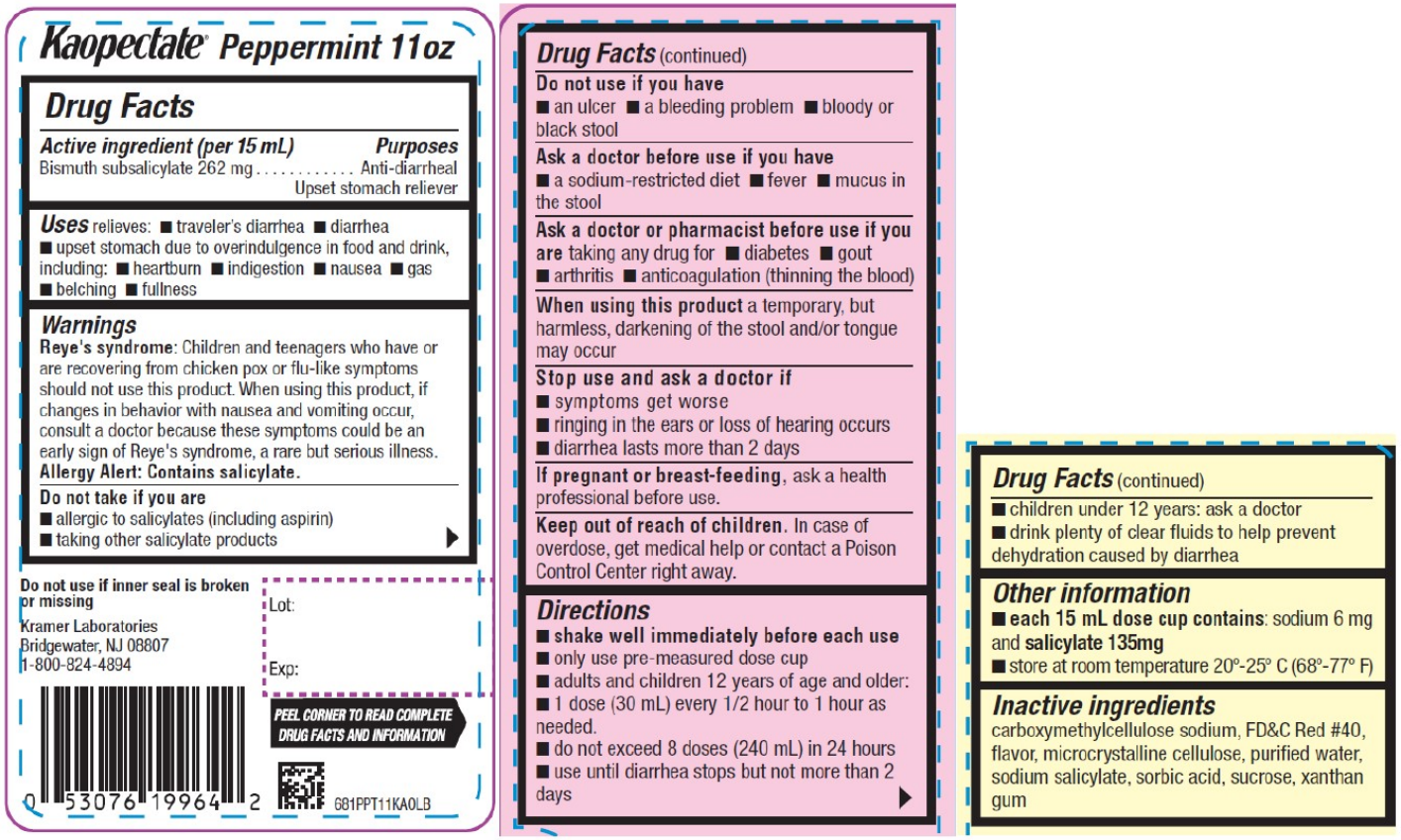
Kramer Laboratories
Bridgewater, NJ 08807
1-800-824-4894

Lot:

Exp:

PEEL CORNER TO READ COMPLETE DRUG FACTS AND INFORMATION

681PPT11KA0LB



Kaopectate® Peppermint 11oz

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Lot:

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PEEL CORNER TO READ COMPLETE DRUG FACTS AND INFORMATION

0 53076 19964 2

681PPT11KA0LB

Drug Facts (continued)

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KAOPECTATE PEPPERMINT FLAVOR ANTI DIARRHEAL

bismuth subsalicylate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55505-199
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (BISMUTH CATION - UNII:ZS9CD1I8YE)	BISMUTH SUBSALICYLATE	262 mg in 15 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Carboxymethylcellulose Sodium, Unspecified (UNII: K679OBS311)	
Fd&C Red No. 40 (UNII: WZB9127XOA)	
Microcrystalline Cellulose (UNII: OP1R32D61U)	
SODIUM SALICYLATE (UNII: WIQ1H85SYP)	
SORBIC ACID (UNII: X045WJ989B)	
SUCROSE (UNII: C151H8M554)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics			
Color		**Score**	
Shape		**Size**	
Flavor	PEPPERMINT	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:55505-199-36	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2020	
2	NDC:55505-199-64	325 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2022	

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