QUALITY CHOICE REGULAR STRENGTH STOMACH RELIEF - bismuth subsalicylate liquid
Chain Drug Marketing Association Inc.

Quality Choice Regular Strength Stomach Relief-527

ACTIVE INGREDIENT(in each 30 mL)

Bismuth subsalicylate 525 mg

PURPOSE

Upset stomach reliever and anti-diarrheal

USE(S)

relieves:

- diarrhea
- heartburn
- indigestion
- nausea
- upset stomach associated with these symptoms

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

- bloody or black stool
- an ulcer
- a bleeding problem

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 of 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 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 immediately.

DIRECTIONS

- **shake well before use**
- mL = milliliter
- TBSP = tablespoon
- adults and children 12 years and over: 1 dose (2 TBSP or 30 mL) every 1/2 to 1 hour as needed
- do not exceed 8 doses (16 TBSP or 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 fluids to help prevent dehydration caused by diarrhea

OTHER INFORMATION

- **each 30 mL or 2 TBSP contains:**
- potassium 25 mg
- salicylate 260 mg

- sodium 8 mg
- protect from freezing
- avoid excessive heat (over 104ºF or 40ºC)
- dosage cup provided

INACTIVE INGREDIENTS

benzoic acid, D&C red # 22, D&C red # 28, flavor, hydroxyethyl cellulose, potassium hydroxide, purified water, saccharin sodium, salicylic acid, simethicone, xanthan gum.

PRINCIPAL DISPLAY PANEL

NDC 83324-020-12

QC QUALITY CHOICE
***Compare to the Active Ingredient in Pepto-Bismol®**

Regular strength
Stomach Relief

Upset Stomach Reliever/Anti-diarrheal
Bismuth Subsalicylate 525 mg per 30 mL

Relieves:

Diarrhea, Heartburn, Indigestion, Nausea & Upset stomach

Original Flavor

12 FL OZ (354 mL)

NDC 83324-020-08

QC QUALITY CHOICE
***Compare to the Active Ingredient in Pepto-Bismol®**

Regular strength
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Upset Stomach Reliever/Anti-diarrheal
Bismuth Subsalicylate 525 mg per 30 mL

Relieves:

Diarrhea, Heartburn, Indigestion, Nausea & Upset stomach

Original Flavor

8 FL OZ (236 mL)

QUALITY CHOICE REGULAR STRENGTH STOMACH RELIEF

bismuth subsalicylate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-020
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII:O414PZ4LPZ)	BISMUTH SUBSALICYLATE	525 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
BENZOIC ACID (UNII: 8SKN0B0MIM)	
D&C RED NO. 22 (UNII: 1678RKX8RT)	
D&C RED NO. 28 (UNII: 767IP0Y5NH)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
HYDROXYETHYL CELLULOSE (1500 MPA.S AT 1%) (UNII: L605B5892V)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	PINK	**Score**	
Shape		**Size**	
Flavor	WINTERGREEN	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-020-12	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/18/2024	
2	NDC:83324-020-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/18/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	part335	04/18/2024	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Establishment

Name	Address	ID/FEI	Business Operations
Guardian Drug Company		119210276	MANUFACTURE(83324-020)

Revised: 4/2024 Chain Drug Marketing Association Inc.