UP AND UP 5 SYMPTOMS DIGESTIVE RELIEF 527 - bismuth subsalicylate 525 mg liquid

Target corporation

up and up 5 symptoms digestive 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

- an ulcer
- bloody or black stool
- 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 104°F or 40°C).
- 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 82442-527-04 compare to active ingredient in Pepto-Bismol®* 5-Symptom Digestive Relief Bismuth Subsalicylate, 525 mg per 30 mL Upset Stomach Reliever/Anti-Diarrheal Relieves upset stomach,

nausea, heartburn, indigestion and diarrhea up&up_{TM} 8 FL OZ (236 mL)



NDC 82442-527-06 compare to active ingredient in Pepto-Bismol®* 5-Symptom Digestive Relief Bismuth Subsalicylate, 525 mg per 30 mL Upset Stomach Reliever/Anti-Diarrheal Relieves upset stomach,

nausea, heartburn, indigestion and diarrhea up&upTM

16 FL OZ (473 mL)



NDC 82442-527-10 compare to active ingredient in Pepto-Bismol®* 5-Symptom Digestive Relief Bismuth Subsalicylate, 525 mg per 30 mL Upset Stomach Reliever/Anti-Diarrheal Relieves upset stomach,

nausea, heartburn,

indigestion and diarrhea

up&upTM

Twin Pack

Two 16 FL OZ (473mL) BOTTLES



UP AND UP 5 SYMPTOMS DIGESTIVE RELIEF 527

bismuth subsalicylate 525 mg liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82442-527
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (BISMUTH CATION - UNII:ZS9CD118YE)	BISMUTH SUBSALICYLATE	525 mg 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)		
POTASSIUM HYDROXIDE (UNII: WZ H3C48M4T)		
WATER (UNII: 059QF0KO0R)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		
SALICYLIC ACID (UNII: O414PZ4LPZ)		
HYDROXYETHYL CELLULOSE (1500 MPA.S AT 1%) (UNII: 1605B5892V)		

DIMETHICONE (UNII: 92RU3N3Y1O)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics			
Color	PINK	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:82442-527- 04	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/18/2024		
2	NDC:82442-527- 06	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/18/2024		
3	NDC:82442-527- 10	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/18/2024		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	part335	06/18/2024	

Labeler - Target corporation (006961700)

Establishment				
Name	Address	ID/FEI	Business Operations	
Guardian Drug Company		119210276	MANUFACTURE(82442-527)	

Revised: 6/2024 Target corporation