UP AND UP DIGESTIVE RELIEF- bismuth subsalicylate suspension
Target Corporation

Target Corporation Digestive Relief Drug Facts

Active ingredient (in each 30 mL dose cup)

Bismuth subsalicylate 525 mg

Purposes

Upset stomach reliever and antidiarrheal

Uses

relieves

- travelers' diarrhea
- diarrhea
- upset stomach due to overindulgence in food and drink, including:
- heartburn
- indigestion
- nausea
- gas
- belching

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

- fever
- mucus in the stool

Ask a doctor or pharmacist before use if you are

taking any drug for

- anticoagulation (thinning 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 a doctor if

- symptoms get worse or last more than 2 days
- 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 (1-800-222-1222).

Directions

- shake well before use
- for accurate dosing, use dose cup
- **adults and children 12 years and over:** 1 dose (30 mL) every 1/2 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 30 mL contains: **sodium 10 mg**
- **salicylate 261 mg**
- low sodium
- sugar free
- store at room temperature
- protect from freezing
- avoid excessive heat (more than over 104°F or 40°C).

Inactive ingredients

carboxymethylcellulose sodium, D&C red #22, D&C red #28, flavor, microcrystalline cellulose, potassium hydroxide, potassium sorbate, purified water, salicylic acid, Simethicone emulsion, sodium benzoate, sucralose, xanthan gum

Questions?

Call 1-800-910-6874

Safety Sealed: DO NOT USE IF IMPRINTED SHRINK BAND IS MISSING OR BROKEN

Principal Display Panel

NDC 11673-706-08

Compare to active ingredient in Pepto-Bismol® Regular Strength*

5-symptom

digestive relief

bismuth subsalicylate 525 mg

upset stomach reliever/antidiarrheal

relieves nausea, heartburn, indigestion, upset stomach and diarrhea

8 FL OZ (236 mL)

Dist. By Target Corp, Mpls , MN 55403

*This product is not manufactured or distributed by Procter & Gamble, distributor of Pepto-Bismol® Regular Strength.

UP AND UP DIGESTIVE RELIEF

bismuth subsalicylate suspension

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-706
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (BISMUTH CATION - UNII:ZS9CD1I8YE, SALICYLIC ACID - UNII:O414PZ4LPZ)	BISMUTH SUBSALICYLATE	525 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
D&C RED NO. 22 (UNII: 1678RKX8RT)	
D&C RED NO. 28 (UNII: 767IP0Y5NH)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
WATER (UNII: 059QF0KO0R)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
DIMETHICONE, UNSPECIFIED (UNII: 92RU3N3Y1O)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics			
Color	PINK (viscous)	**Score**	
Shape		**Size**	
Flavor	WINTERGREEN	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-706-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/15/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M008	03/15/2018	

Labeler - Target Corporation (006961700)

Revised: 11/2024 Target Corporation