HEB STOMACH RELIEF MAX STRENGTH- bismuth subsalicylate suspension HEB

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

HEB Max Strength Bismuth Subsalicylate Drug Facts

Active ingredient (in each 30 mL dose cup)

Bismuth subsalicylate 1050 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 at (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 hour as needed
- do not exceed 4 doses (120 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 20 mg
- salicylate 437 mg
- low sodium
- sugar free
- store at room temperature.

- protect from freezing
- avoid excessive heat (more than 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 or comments?

1-866-467-2748

Principal Display Panel

Compare to Pepto-Bismol® Maximum Strength active ingredient

HEB_®

Stomach Relief

Max

Bismuth Subsalicylate 1050 mg

Upset Stomach Reliever/Antidiarrheal

Multi-Symptom Relief

Relieves:

- Nausea
- Heartburn
- Indigestion
- Upset Stomach
- Diarrhea

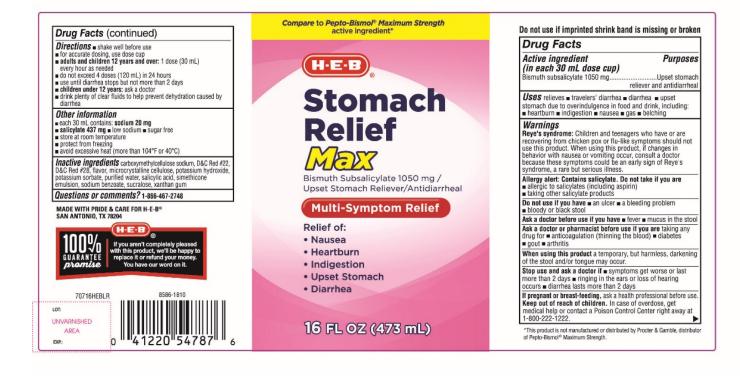
MADE WITH PRIDE & CARE FOR H-E-B®

SAN ANTONIA, TX 78204

100% GUARANTEE promise, if you aren't completely pleased with this product, we'll be happy to replace it or refund your money. You have our word on it.

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

Package Label 473 mL



Package Label 236 mL



HEB STOMACH RELIEF MAX STRENGTH

bismuth subsalicylate suspension

Product Information

| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:37808-770 |
|-------------------------|----------------|--------------------|---------------|
| Route of Administration | ORAL | | |

| Active Ingredient/Active Moiety | | |
|--|--------------------------|---------------------|
| Ingredient Name | Basis of Strength | Strength |
| BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII:O414PZ4LPZ, BISMUTH CATION - UNII:ZS9CD118YE) | BISMUTH SUBSALICYLATE | 1050 mg in 30 mL |

| Inactive Ingredients | | |
|---|----------|--|
| Ingredient Name | Strength | |
| CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311) | | |
| 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 (UNII: 92RU3N3Y1O) | | |
| SODIUM BENZOATE (UNII: OJ245FE5EU) | | |
| SUCRALOSE (UNII: 96K6UQ3ZD4) | | |
| XANTHAN GUM (UNII: TTV12P4NEE) | | |

| Product Characteristics | | | |
|-------------------------|----------------|--------------|--|
| Color | PINK (viscous) | Score | |
| Shape | | Size | |
| Flavor | | Imprint Code | |
| Contains | | | |

| ı | P | Packaging | | | |
|---|---|----------------------|---|-------------------------|-----------------------|
| | # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| | 1 | NDC:37808-770- 16 | 473 mL in 1 BOTTLE; Type 0: Not a Combination Product | 04/23/2019 | |
| | 2 | NDC:37808-770- 08 | 236 mL in 1 BOTTLE; Type 0: Not a Combination Product | 04/23/2019 | |

| Marketing Information | | | |
|-----------------------|---|-------------------------|-----------------------|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC monograph final | part335 | 04/23/2019 | |
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Labeler - HEB (007924756)

Revised: 9/2023 HEB