ANTI-DIARRHEAL MAXIMUM STRENGTH- bismuth subsalicylate tablet
Wal-Mart Stores Inc

Equate 44-749

Active ingredient (in each caplet)

Bismuth subsalicylate 525 mg

Purpose

Antidiarrheal

Uses

relieves
- travelers' diarrhea
- diarrhea

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)
- gout
- diabetes
- 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
- 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

- drink plenty of clear fluids to help prevent dehydration caused by diarrhea
- swallow with water; do not chew
- adults and children 12 years and over: 1 caplet every 1/2 hour or 2 caplets every hour as needed
- do not exceed 8 caplets in 24 hours
- use until diarrhea stops but not more than 2 days
- children under 12 years: ask a doctor

Other information

- **each caplet contains:** calcium 45 mg, salicylate 206 mg, sodium 3 mg
- store at 25ºC (77ºF); excursions permitted between 15º-30ºC (59º-86ºF)
- avoid excessive heat
- use by expiration date on package

Inactive ingredients

calcium carbonate, corn starch, D&C red #27 aluminum lake, D&C red #30 aluminum lake, magnesium stearate, mannitol, microcrystalline cellulose, povidone, silicon dioxide, sodium starch glycolate, stearic acid

Questions or comments?

1-888-287-1915

Principal display panel

equate™

NDC 49035-769-08

Compare to the active ingredient in Pepto® Diarrhea*

MAXIMUM STRENGTH
Anti-Diarrheal
Bismuth Subsalicylate 525 mg
Anti-Diarrheal
Relieves Diarrhea

Actual Size

525 mg EACH

24 CAPLETS

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716

*This product is not manufactured or distributed by The Procter & Gamble Company, owner of the registered trademark Pepto® Diarrhea.

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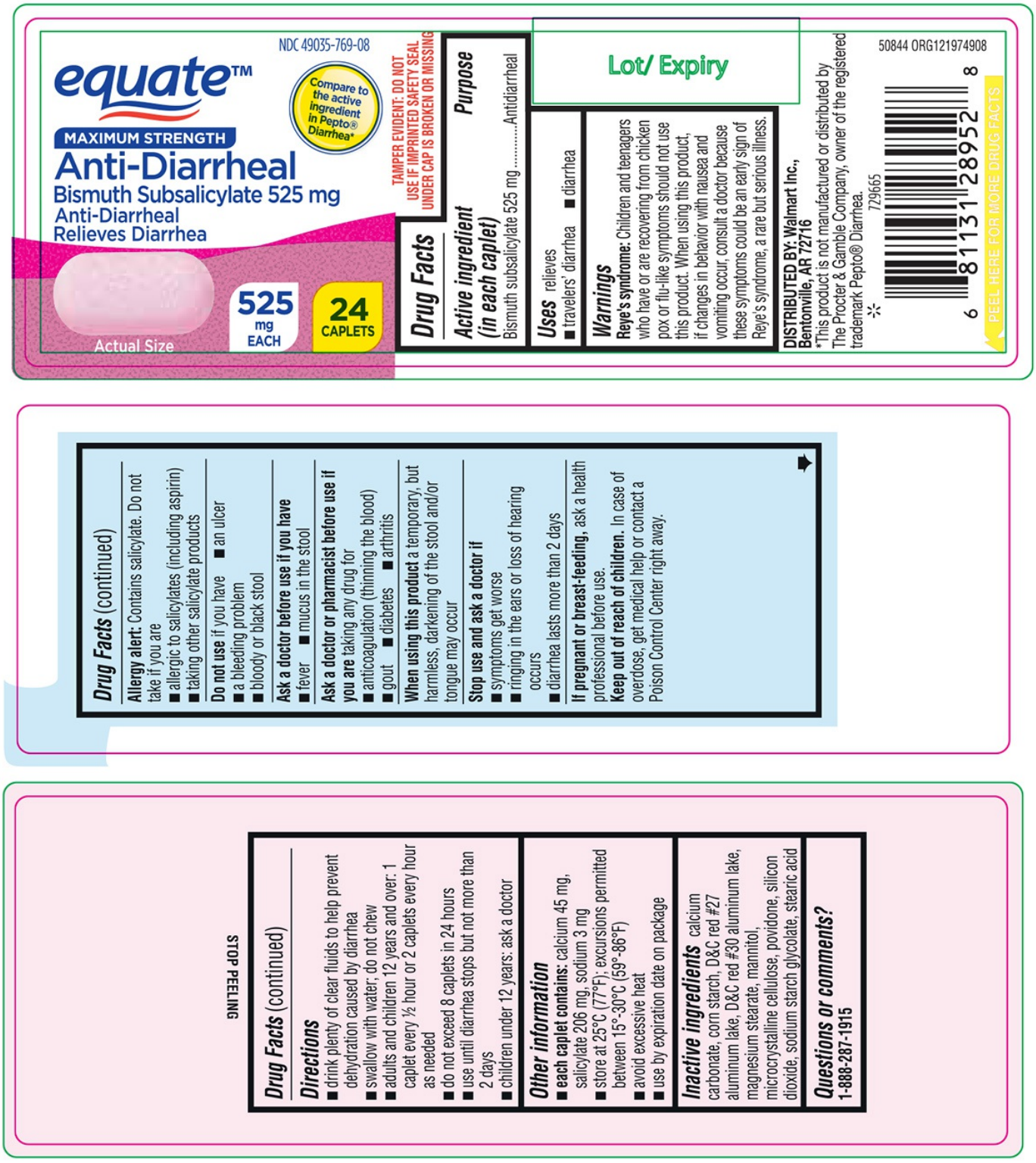

Equate 44-749

ANTI-DIARRHEAL MAXIMUM STRENGTH

bismuth subsalicylate tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-769

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:ZS9CD1I8YE)	BISMUTH SUBSALICYLATE	525 mg

Inactive Ingredients

Ingredient Name	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK)	
STARCH, CORN (UNII: O8232NY3SJ)	
D&C RED NO. 27 ALUMINUM LAKE (UNII: ZK64F7XSTX)	
D&C RED NO. 30 ALUMINUM LAKE (UNII: GE75M6YV5W)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MANNITOL (UNII: 3OWL53L36A)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	pink	**Score**	no score
Shape	OVAL	**Size**	19mm
Flavor		**Imprint Code**	44;749
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-769-08	24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/06/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M008	04/06/2020	

Labeler - Wal-Mart Stores Inc (051957769)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(49035-769)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(49035-769)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(49035-769)

Revised: 3/2025

Wal-Mart Stores Inc