MEDIQUE DIOTAME- bismuth subsalicylate tablet, chewable
Unifirst First Aid Corporation

Medique Diotame

Drug Facts

Active ingredient (in each tablet)

Bismuth Subsalicylate 262 mg

(each tablet contains 102 mg salicylate)

Purpose

antidiarrheal/antacid

Uses

relieves
- traveler's diarrhea
- diarrhea
- upset stomach reliever due to overindulgence in food and drink, including: • heartburn • indigestion • nausea • gas • belching • fullness

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 if you are taking any drug for
- anticoagulation (thinning of the blood)
- diabetes
- gout
- arthritis

When using this product a temporary and harmless darkening of the tongue and/or

stool may occur.

Stop use and ask a doctor if

- symptoms get worse or lasts 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 accidental overdose, contact a physician or poison control center immediately (1-800-222-1222).

Directions

- chew or dissolve in mouth
- do not swallow tablets whole
- drink plenty of clear fluids to help prevent dehydration, which may accompany diarrhea

Adults and children: (12 years and over)

- chew 2 tablets every 1/2 to 1 hour as needed
- do not exceed 16 tablets in 24 hours
- use until diarrhea stops but not more than 2 days

Children under 12 years:

ask a doctor

Other information

- **phenylketonurics:** contains phenylalanine 1.1mg per tablet
- **calcium** content per tablet: 73 mg
- store at room temperature 59°-86°F (15°-30°C)
- tamper-evident sealed packets
- do not use any opened or torn packets

Inactive ingredients

acacia gum, aspartame, calcium carbonate, D&C red #27 aluminum lake, dextrates, flavoring, magnesium stearate, maltodextrin, microcrystalline cellulose, peppermint flavor, silicon dioxide

Questions or comments? 1-800-634-7680

Medique Diotame Label

Medique®

Diotame

Bismuth Subsalicylate

Chewable Tablets

Upset Stomach/Diarrhea • Bismuth Subsalicylate 262 mg

Tamper Evident Unit Dose Packets

MEDIQUE DIOTAME

bismuth subsalicylate tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47682-210
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	262 mg

Inactive Ingredients

Ingredient Name	Strength
ASPARTAME (UNII: Z0H242BBR1)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
DEXTRATES (UNII: G263MI44RU)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
ACACIA (UNII: 5C5403N26O)	

Product Characteristics			
Color	pink	**Score**	no score
Shape	ROUND	**Size**	16mm
Flavor	PEPPERMINT	**Imprint Code**	RH;046
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-210-83	15 in 1 BOX	04/01/2014	
1	NDC:47682-210-99	2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:47682-210-33	50 in 1 BOX	04/01/2014	
2		2 in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:47682-210-13	250 in 1 BOX	04/01/2014	
3		2 in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:47682-210-64	12 in 1 BOX	04/01/2014	
4		2 in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:47682-210-99	2 in 1 PACKET; Type 0: Not a Combination Product	04/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M008	04/01/2014	

Labeler - Unifirst First Aid Corporation (832947092)

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