MI ACID- aluminum hydroxide, magnesium hydroxide, simethicone suspension ATLANTIC BIOLOGICALS CORP.

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.

Major Pharmaceuticals Mi-Acid™ Drug Facts

Active ingredients (in each 10 mL)

Aluminum hydroxide (equiv. to dried gel, USP) 400 mg Magnesium hydroxide 400 mg Simethicone 40 mg

Purposes

Antacid

Antigas

Uses

relieves

- heartburn
- acid indigestion
- sour stomach
- upset stomach due to these symptoms
- pressure and bloating commonly referred to as gas

Warnings

Ask a doctor before use if you have

- kidney disease
- a magnesium-restricted diet

Ask a doctor or pharmacist before use if you are

presently taking a prescription drug. Antacids may interact with certain prescription drugs.

When using this product

do not exceed 120 mL in a 24-hour period, or use the maximum dosage for more than 2 weeks, unless directed by a doctor

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
- measure with dosing cup provided
- **adults and children 12 years and over:** 10 mL 20 mL between meals, at bedtime or as directed by a doctor
- do not take more than 120 mL in any 24 hour period
- do not use the maximum dosage for more than 2 weeks
- children under 12 years: ask a doctor
- mL = milliliter

Other information

- each 10 mL contains: magnesium 175 mg
- does not meet USP requirements for preservative effectiveness
- store at 20-25°C (68-77°F), do not freeze

Inactive ingredients

butylparaben, flavor, hypromellose, microcrystalline cellulose and carboxymethylcellulose sodium, peppermint oil, propylparaben, purified water, simethicone emulsion, sorbitol, sorbitol solution

Questions or comments?

1-800-616-2471

Principal Display Panel

FAST ACTING

MI-ACIDTM

Regular Strength

ANTACID / ANTIGAS

CLASSIC

SOOTHING RELIEF OF:

HEARTBURN

ACID INDIGESTION

SOUR STOMACH

ALCOHOL 0.2%

17856-6833-01

Aluminum

Hydroxide/Magnesium

Hydroxide

Simethicone (MI-ACID)

Delivers 30mL

See package insert for indications and dosage schedule

ANTACIDI ANTIGAS, ALCOHOL 0.2%. Store at 20°-25°(68°-77°F). Do not freeze. KEEP THIS AND ALL DRUGS OUT OF THE REACH OF CHILDREN



17856-6833-01

Aluminum

Hydroxide/Magnesium

Hydroxide/Simethicone

(MI-ACID)

GTIN: 00117856683315

S/N: 01173201

Exp: 03/22/21

Lot: 011732

OTC

Packaged by:Unit Dose Solutions Morrisville, NC 27560 Distributed by: AtlanticBiologicals Corp, Miami Fl 33179

Dosage: 1200mg/1200mg/120 mg /30mL

Qty: 50 CUPS

Rev.09/19

Call to Reorder: 800.509.7592

MI ACID

aluminum hydroxide, magnesium hydroxide, simethicone suspension

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:17856-6833(NDC:0904-6833)	
Route of Administration	ORAL			

l	Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength	
	AT TIMINTIM HVDDA YIDE (TIMII- SARATSHIMA) (AT TIMINTIM HVDDA YIDE _ TIMI-SARATSHIMA)	ALUMINUM	400 mg	

ALCIVILIONI III DAGAIDE (CIVIL. SQDU IZICINU) (ALCIVILIONIVI III DAGAIDE - CIVIL.SQDU IZICINU)	HYDRO XIDE	in 10 mL
MAGNESIUM HYDRO XIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6 V3LHY838, HYDRO XIDE ION - UNII:9159 UV381P)	MAGNESIUM HYDRO XIDE	400 mg in 10 mL
DIMETHICO NE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	40 mg in 10 mL

Inactive Ingredients				
Ingredient Name	Strength			
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)				
BUTYLPARABEN (UNII: 3QPI1U3FV8)				
HYPROMELLOSES (UNII: 3NXW29V3WO)				
CELLULOSE, MICRO CRYSTALLINE (UNII: OP1R32D61U)				
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679 OBS 311)				
PEPPERMINT OIL (UNII: AV092KU4JH)				
PROPYLPARABEN (UNII: Z8 IX2SC1OH)				
WATER (UNII: 059QF0KO0R)				
SORBITOL (UNII: 506T60A25R)				

Product Characteristics					
Color	WHITE (opaque)	Score			
Shape		Size			
Flavor	MINT, LEMON	Imprint Code			
Contains					

F	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:17856-6833- 1	50 in 1 BOX, UNIT-DOSE	09/23/2020			
1		30 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part331	12/28/2018		

Labeler - ATLANTIC BIOLOGICALS CORP. (047437707)

Establishment			
Name	Address	ID/FEI	Business Operations
ATLANTIC BIOLOGICALS CORP.		047437707	relabel(17856-6833), repack(17856-6833)

Revised: 9/2020 ATLANTIC BIOLOGICALS CORP.