SODIUM BICARBONATE ANTACID- sodium bicarbonate tablet Rugby Laboratories

4621 - Rugby

Drug Facts

Active ingredient (in each tablet)

Sodium bicarbonate 5 gr (325 mg)

Purpose

Antacid

Uses

relieves

- acid indigestion
- heartburn
- sour stomach
- upset stomach associated with these symptoms

Warnings

Ask a doctor before use if you have a sodium restricted diet

Ask a doctor or pharmacist before use if you are taking a prescription drug.

Antacids may interact with certain prescription drugs.

Stop use and ask a doctor if symptoms last more than 2 weeks

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

Directions

- do not use the maximum dosage for more than 2 weeks
- tablets may be swallowed whole or dissolved in water prior to use
- adults 60 years of age and over: 2-4 tablets every 4 hours, not more than 24 tablets in 24 hours
- adults under 60 years of age: 2-8 tablets every 4 hours, not more than 48 tablets in 24 hours

Other information

• each tablet contains: sodium 89 mg

• Store in a dry place at 15° - 30°C (59° - 86° F)

Inactive ingredients

croscarmellose sodium, microcrystalline cellulose, silica, stearic acid, and talc

Questions or comments?

1-800-231-4670

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

WARNING: This product can expose you to chemicals including lead and mercury which are known to the State of

California to cause birth defects or other reproductive harm. For more information go to www.P65Warnings.ca.gov.

DISTRIBUTED BY:

RUGBY® LABORATORIES

INDIANAPOLIS, IN 46268

Questions or comments?

Call (800) 616-2471

www.majorpharmaceuticals.com

Rev. 10/23 R-29

Rugby

NDC 0536-1046-10

Sodium Bicarbonate 5 gr (325 mg)

ANTACID

1000 WHITE TABLETS



NDC 0536-1046-10

Sodium **Bicarbonate 5 gr** (325 mg)

ANTACID

1000 WHITE TABLETS

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

Purpose Antacid

Active ingredient (in each tablet)

Sodium bicarbonate 5 gr (325 mg)

Uses relieves • acid indigestion • heartbum • sour stomach • upset stomach associated with these

Ask a doctor or pharmacist before use if you are taking a prescription drug. Ask a doctor before use if you have a sodium restricted diet

Antacids may interact with certain prescription drugs

Stop use and ask a doctor if symptoms last more than 2 weeks

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

Directions

tablets may be swallowed whole or dissolved in water prior to use do not use the maximum dosage for more than 2 weeks

adults 60 years of age and over: 2-4 tablets every 4 hours, not more tran 24 tablets in 24 hours
adults under 60 years of age: 2-8 tablets every 4 hours, not more tran 48 tablets in 24 hours

• each tablet contains; sodium 89 mg • Store in a dry place at 15° – 30°C (59° – 86°F). Other information

proscamellose sodium, microcrystalline cellulose, silica, stearic acid and talc nactive ingredients

Questions or comments?

Questions or comme Call (800) 616-2471 /Exp.Date:

SODIUM BICARBONATE ANTACID

sodium bicarbonate tablet

Product Information

Product Type HUMAN OTC DRUG NDC:0536-1046 Item Code (Source)

Route of Administration ORAL

Active Ingredient/Active Moiety Basis of Ingredient Name Strength Strength **SODIUM** SODIUM BICARBONATE (UNII: 8MDF5V39QO) (SODIUM CATION -325 mg **BICARBONATE** UNII:LYR4M0NH37)

Inactive Ingredients Strength **Ingredient Name** CROSCARMELLOSE SODIUM (UNII: M280L1HH48) CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U) STEARIC ACID (UNII: 4ELV7Z65AP) SILICON DIOXIDE (UNII: ETJ7Z6XBU4) TALC (UNII: 7SEV7J4R1U)

Product Characteristics

Color white Score no score

Shape	ROUND	Size	10mm
Flavor		Imprint Code	CPC76
Contains			

l	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:0536-1046- 10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	03/25/2015		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M001	03/25/2015			

Labeler - Rugby Laboratories (079246066)

Revised: 12/2023 Rugby Laboratories