SODIUM BICARBONATE- sodium bicarbonate tablet
Richmond Pharmaceuticals, Inc.

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.

SODIUM BICARBONATE 10 gr (650 mg)

Drug Facts

Active Ingredient (in each tablet)

Sodium bicarbonate 10 gr (650 mg)

PURPOSE

Antacid

Indications:

Relieves:

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

Warnings

Ask a doctor or pharmacist

- if you are on a sodium-restricted diet.
- if you are taking a prescription drug. Antacids may interact with certain prescription drugs.
- if symptons last more than 2 weeks

As with any drug, if you are pregnant or nursing a baby, seek advise of a health professional before using this product.

Directions:

- Adults 60years of age and over - 1-2 tablets every 4 hours. Not more than 12 tablets in 24 hours
- Adults under 60 years- 1-4 tablets every4 hours. Not more than 24 tablets in 24 hours
- Dissolve tabelt completely in water before drinking.
- DO NOT EXCEED RECOMMENDED DOSE. Not recommended for children.

Other Information:

- each tablet contains: sodium 178 mg
- store at room temperature 15 °- 30 °C (59 °- 86 °F).

KEEP THIS AND ALL DRUGS OUT OF THE REACH OF CHILDREN.

In case of accidental overdose, seek professional assistance or contact a poison control center

immediately.

INACTIVE INGREDIENT

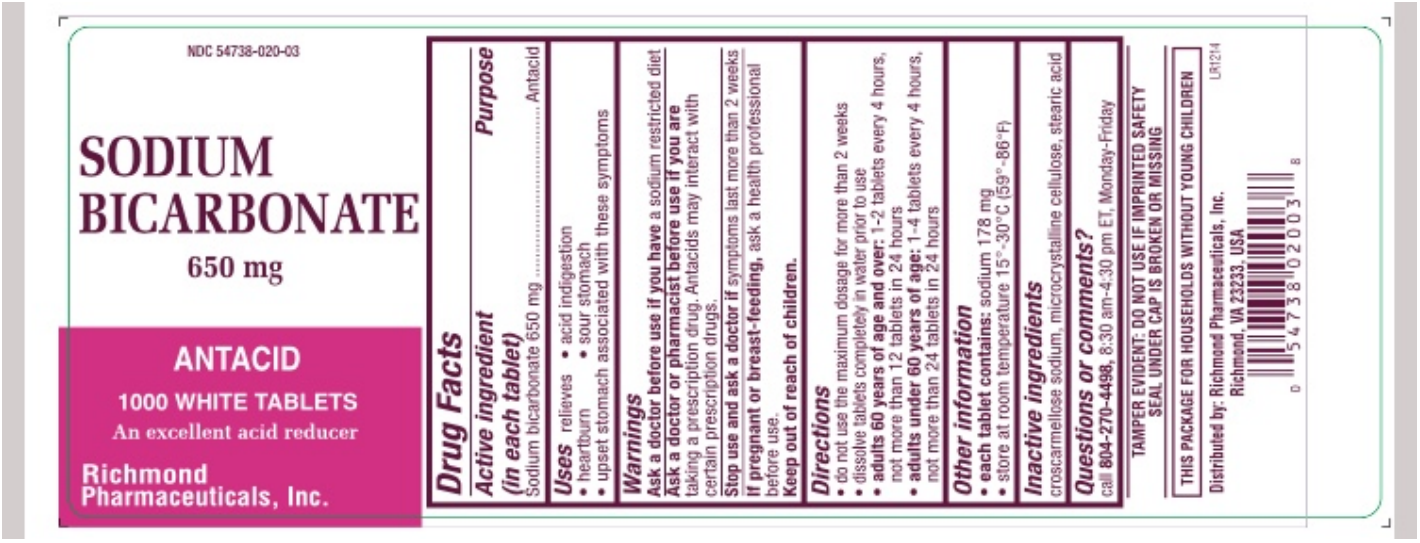
croscarmellose sodium, microcrystalline cellulose, stearic acid

Questions or Comments

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

call 804-270-4498, 8.30 am-4.30 pm ET, Monday - Friday

Principal display panel



SODIUM BICARBONATE

sodium bicarbonate tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54738-020
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM BICARBONATE (UNII: 8MDF5V39QO) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM BICARBONATE	650 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics			
Color	white (White)	**Score**	no score
Shape	ROUND (round)	**Size**	11mm
Flavor		**Imprint Code**	AP;119
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54738-020-03	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	05/01/2015	

Labeler - Richmond Pharmaceuticals, Inc. (043569607)

Registrant - Advance Pharmaceutical Inc. (078301063)

Establishment

Name	Address	ID/FEI	Business Operations
Advance Pharmaceutical Inc.		078301063	manufacture(54738-020)

Revised: 10/2017 Richmond Pharmaceuticals, Inc.