

**SODIUM BICARBONATE 10 GR- sodium bicarbonate tablet**  
**GRAXCELL PHARMACEUTICAL, LLC**

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**Sodium Bicarbonate 10 gr (650 mg), 70795-1190**

**Active ingredient (in each tablet)**

Sodium bicarbonate 10 gr (650 mg)

**Purpose**

Antacid

**Uses**

Relieves

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

**Warnings**

**Do not use**

- this product if you are on a sodium-restricted diet unless directed by a doctor
- Do not take more than 24 tablets for adults up to 60 years of age (or 12 tablets for adults 60 years of age or older) in a 24 hour period nor use maximum dosage for more than 2 weeks, except under the advice and supervision of a physician.
- As with any drug, if you are pregnant or nursing a baby, seek advice of a health professional before using this product

Drug Interaction Precaution

**Stomach Warning**

- **TO AVOID SERIOUS INJURY, DO NOT TAKE UNTIL TABLET IS COMPLETELY DISSOLVED IT IS VERY IMPORTANT NOT TO TAKE THIS PRODUCT WHEN OVERLY FULL FROM FOOD OR DRINK**
- Consult a doctor if severe stomach pain occurs after taking this product.

**Drug Interaction Precaution**

- Ask a physician or pharmacist before use if you are presently taking a prescription drug
- **Antacids may interact with certain prescription drugs**

**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

## Directions

**Adults:** Take 1 tablet, dissolved in a glass of water, as needed. Maximum daily dose for adults up to 60 years of age is 24 tablets.

Maximum daily dose for adults 60 years of age or older is 12 tablets. Dissolve completely in water before drinking.

## DO NOT EXCEED RECOMMENDED DOSE

Not recommended for children.

## Other information

- each tablet contains : **sodium 178 mg (7.74 meq)**
- store at room temperature 15°-30°C (59°-86°F) in well-closed containers as defined in the USP

## Inactive ingredients

Pregelatinized starch, NF and mineral oil, USP

## Questions or comments?

Call toll free 1-888-266-8818

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

**GRAXCELL**  
PHARMACEUTICAL, LLC.

# Sodium Bicarbonate

Sodium Bicarbonate 10 gr (650 mg)

## Antacid

# 1000 Tablets

Drug Facts	
<b>Active ingredient (in each tablet)</b>	<b>Purpose</b>
Sodium Bicarbonate 10 gr (650 mg) .....	Antacid
<b>Uses</b>	
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<b>Warnings</b>	
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<b>Stomach Warning: TO AVOID SERIOUS INJURY, DO NOT TAKE UNTIL TABLET IS COMPLETELY DISSOLVED IT IS VERY IMPORTANT NOT TO TAKE THIS PRODUCT WHEN OVERLY FULL FROM FOOD OR DRINK.</b> Consult a doctor if severe stomach pain occurs after taking this product.	
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<b>DO NOT EXCEED RECOMMENDED DOSE</b> Not recommended for children.	
<b>Other information</b>	
■ each tablet contains: sodium 178 mg (7.74 meq) ■ store at room temperature 15°-30°C (59°-86°F) in well-closed containers as defined in the USP	
<b>Inactive ingredients</b> Pregelatinized starch, NF and mineral oil, USP	
<b>Questions or comments?</b> Call toll free 1-888-266-8818	

Manufactured by:  
GRAXCELL Pharmaceutical LLC  
136 OAK Drive  
Syosset, N.Y. 11791

Distributed by:  
GRAXCELL Pharmaceutical LLC  
130 Knickerbocker Ave  
Bohemia, NY 11716  
MADE IN USA



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## SODIUM BICARBONATE 10 GR

sodium bicarbonate tablet

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:70795-1190
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>SODIUM BICARBONATE</b> (UNII: 8MDF5V39QO) (SODIUM CATION - UNII:LYR4MONH37)	SODIUM BICARBONATE	650 mg

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>STARCH, CORN</b> (UNII: O8232NY35J)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	2 pieces
<b>Shape</b>	ROUND	<b>Size</b>	11mm
<b>Flavor</b>		<b>Imprint Code</b>	G35
<b>Contains</b>			

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:70795-1190-0	1000 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2017	
2	NDC:70795-1190-5	500 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2017	
3	NDC:70795-1190-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2017	
4	NDC:70795-1190-2	200 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2017	

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M001	10/02/2017	

**Labeler** - GRAXCELL PHARMACEUTICAL, LLC (056556923)**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
GRAXCELL PHARMACEUTICAL, LLC		056556923	manufacture(70795-1190)

