PREMIER VALUE SODIUM BICARBONATE- sodium bicarbonate powder
Chain Drug Consortium

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.

Premier Value Sodium Bicarbonate

Drug Facts

Active Ingredient

Soduim Bicarbonate, USP

Purpose

Antacid

Use

For relief of heartburn, acid indigestion, and upset stomach associated with these symptoms.

Warnings
NOT FOR INJECTIONS

Except under supervision of a doctor do not administer to children under 6 years of age.

Do not take more than six, 1/2 tsp. per person up to 60 years old, or three 1/2 tsp. per person 60 years or older in a 24 hour period.

Do not use this product if you are on a sodium restricted diet (each 1/2 tsp. contains 30 mEq (0.7 g) Sodium).

Do not use the maximum does more than 2 weeks.

Ask a doctor or pharmacist before use if

you are currently taking a prescription drug. Antacids may interact with certain prescription drugs.

Keep out of reach of children.

In case of accidental overdose, seek professional assistance or contact a Poison Control Center immediately.

Directions

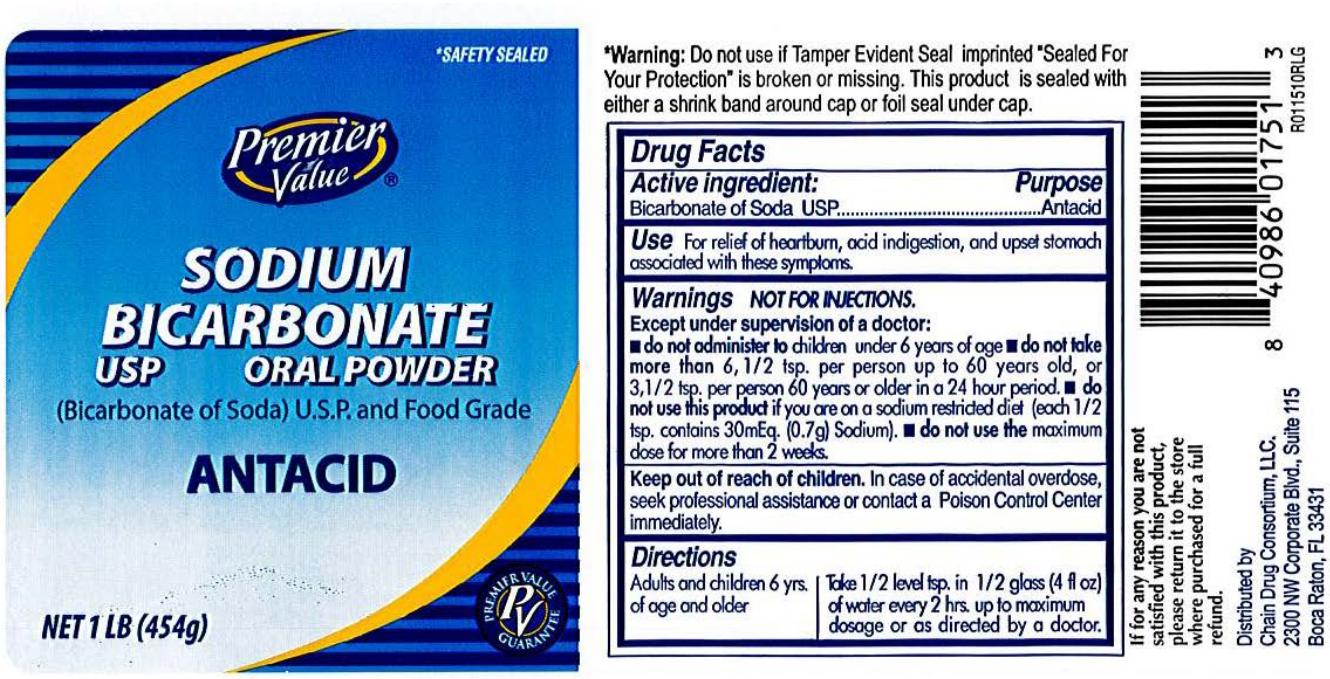
Adult and children 6 yrs of age and older:

Take 1/2 tsp. in 1/2 g;ass (4 fl oz) of water every 2 hrs. up to maximum dosage or as directed by doctor.

Inactive ingredients

none

Label



PREMIER VALUE SODIUM BICARBONATE

sodium bicarbonate powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68016-685
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM BICARBONATE (UNII: 8MDF5V39QO) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM BICARBONATE	1000 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:68016-685-01	454 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/27/2017	
2	NDC:68016-685-94	113 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/27/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	10/27/2017	

Labeler - Chain Drug Consortium (101668460)

Registrant - Humco Holding Group, Inc. (825672884)

Establishment

Name	Address	ID/FEI	Business Operations
Humco Holding Group, Inc.		825672884	label(68016-685) , manufacture(68016-685) , pack(68016-685) , analysis(68016-685)

Revised: 10/2017 Chain Drug Consortium