HEARTBURN RELIEF MAXIMUM STRENGTH- ranitidine tablet, film coated
GREENBRIER INTERNATIONAL, INC.

Assured 44-620-Delisted

Active ingredient
(in each tablet)

Ranitidine 150 mg
(as ranitidine hydrochloride USP, 168 mg)

Purpose

Acid reducer

Uses

- relieves heartburn associated with acid indigestion and sour stomach
- prevents heartburn associated with acid indigestion and sour stomach brought on by certain foods and beverages

Warnings

Allergy alert: Do not use if you are allergic to ranitidine or any other acid reducers.

Do not use

- if you have trouble or pain swallowing food, vomiting with blood, or bloody or black stools. These may be signs of a serious condition. See your doctor.
- with other acid reducers
- if you have kidney disease, except under the advice and supervision of a doctor

Ask a doctor before use if you have

- frequent **chest pain**
- frequent wheezing, particularly with heartburn
- unexplained weight loss
- nausea or vomiting
- stomach pain
- had heartburn over 3 months. This may be a sign of a more serious condition.
- heartburn with **lightheadedness, sweating, or dizziness**
- chest pain or shoulder pain with shortness of breath; sweating; pain spreading to arms, neck or shoulders; or lightheadedness

Stop use and ask a doctor if

- your heartburn continues or worsens
- you need to take this product for more than 14 days

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- adults and children 12 years and over:
  - to **relieve symptoms**, swallow 1 tablet with a glass of water
  - to **prevent symptoms**, swallow 1 tablet with a glass of water **30 to 60 minutes before** eating food or drinking beverages that cause heartburn
  - can be used up to twice daily (do not take more than 2 tablets in 24 hours)
- children under 12 years: ask a doctor

Other information

- **TAMPER EVIDENT: DO NOT USE IF THE CARTON OR PRINTED FOIL UNDER CAP IS OPEN OR TORN**
- store at 20º-25ºC (68º-77ºF)
- avoid excessive heat or humidity
- this product is sodium and sugar free
- see end flap for expiration date and lot number

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide red, magnesium stearate, microcrystalline cellulose, polyethylene glycol, talc, titanium dioxide

Questions or comments?

Call 1-800-426-9391 8:30 AM-4:00 PM ET, Monday-Friday

Principal display panel

ASSURED™

COMPARE TO ACTIVE INGREDIENT OF **MAXIMUM STRENGTH ZANTAC 150®***

MAXIMUM STRENGTH
HEARTBURN
RELIEF
• **Ranitidine 150 mg** - *Acid Reducer*

Relieves and prevents heartburn due to acid indigestion

30 tablets

TAMPER EVIDENT: DO NOT USE IF THE CARTON OR PRINTED FOIL UNDER CAP IS OPEN OR TORN

DOES NOT CONTAIN GLUTEN
*This product is not manufactured or distributed by Boehringer Ingelheim Pharmaceuticals, Inc., owner of the registered trademark Zantac 150®.
50844 ORG031362001
Product of India

ITEM #200907
Distributed by: **Greenbrier International, Inc.**
Chesapeake, VA 23320 USA

Assured 44-620

HEARTBURN RELIEF MAXIMUM STRENGTH

ranitidine tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:33992-0620
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
RANITIDINE HYDROCHLORIDE (UNII: BK76465IHM) (RANITIDINE - UNII:884KT10YB7)	RANITIDINE	150 mg

Inactive Ingredients	
Ingredient Name	**Strength**
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	PINK	**Score**	no score
Shape	OVAL	**Size**	12mm
Flavor		**Imprint Code**	9R
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:33992-0620-1	1 in 1 CARTON	06/15/2013	07/15/2018
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA200536	06/15/2013	07/15/2018

Labeler - GREENBRIER INTERNATIONAL, INC. (610322518)

Revised: 4/2016 GREENBRIER INTERNATIONAL, INC.