RANITIDINE- ranitidine tablet
Dr. Reddy's Laboratories Limited

Drug Facts

Active ingredient (in each tablet)

Ranitidine 75 mg (as ranitidine hydrochloride, USP 84 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 eating or drinking certain foods and beverages

Warnings

Allergy alert: Do not use if you are allergic to ranitidine or 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

Ask a doctor before use if you have

- 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
- frequent **chest pain**
- frequent wheezing, particularly with heartburn
- unexplained weight loss
- nausea or vomiting
- stomach pain
- kidney disease

Ask a doctor or pharmacist before use if you are

• taking a prescription drug. Acid reducers may interact with certain prescription drugs.

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 right away (1-800-222-1222).

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)
  - do not chew tablet
- children under 12 years: ask a doctor

Other information

- this product is sodium and sugar free
- Blister: do not use if individual blister unit is open or torn
  Bottle: do not use if printed foil under bottle cap is open or torn
- avoid excessive heat or humidity
- store at 20°-25°C (68°-77°F)

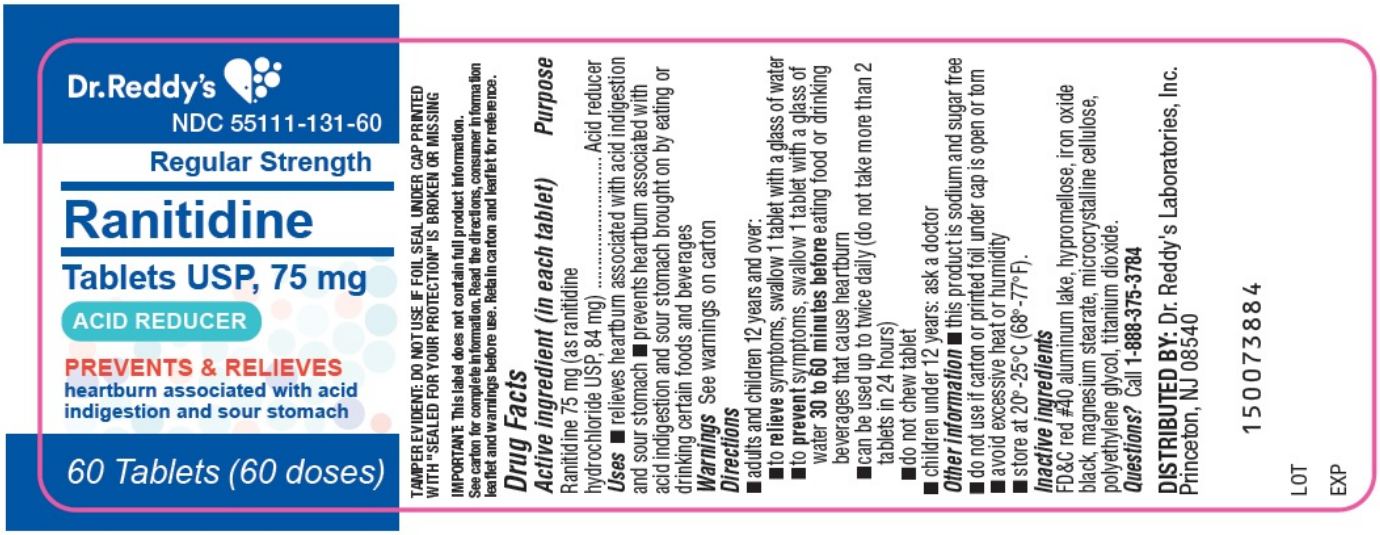
Inactive ingredients

FD&C red #40 aluminum lake, hypromellose, iron oxide black, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide.

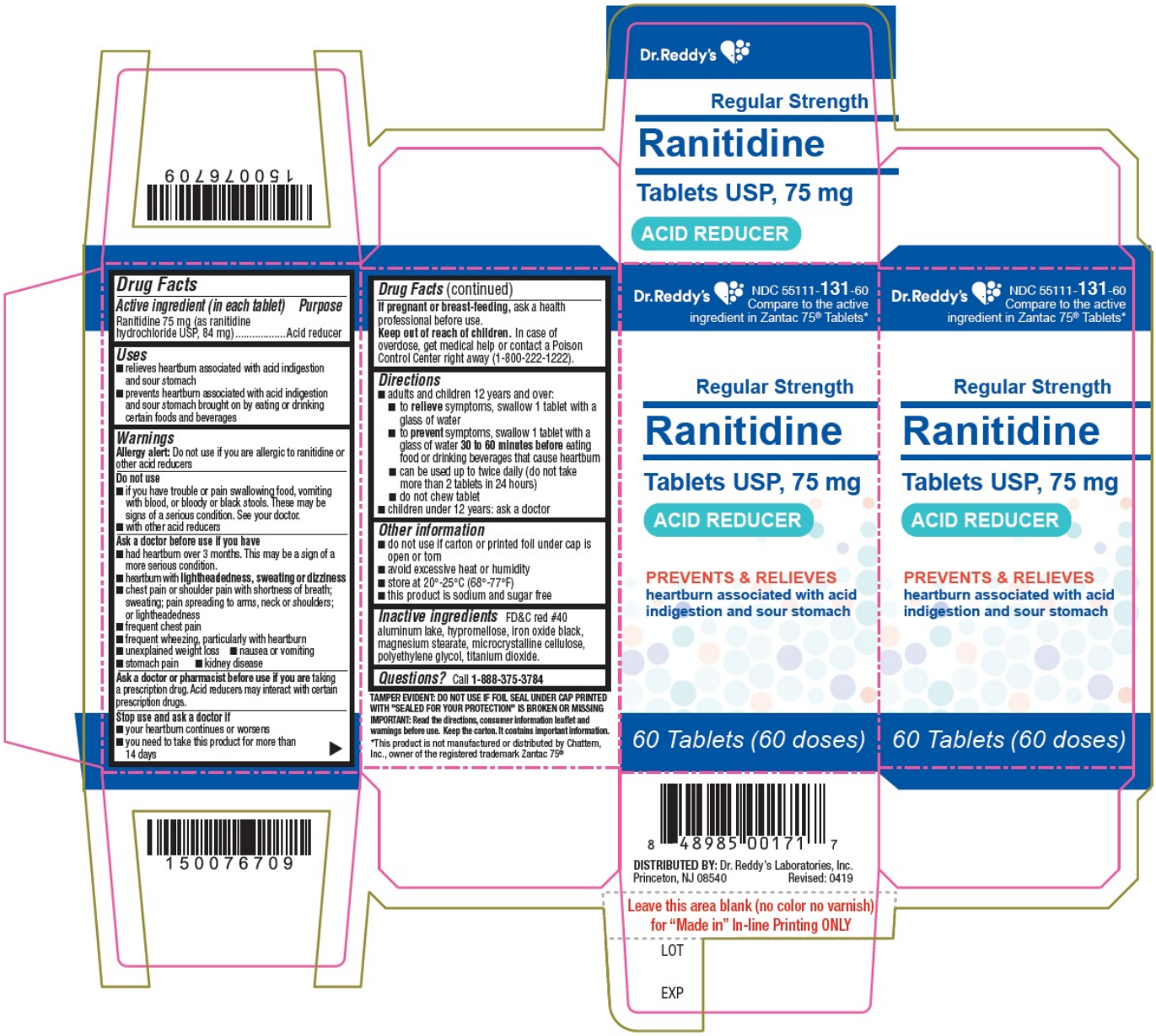
Questions? call **1-888-375-3784**

Read the directions, consumer information leaflet and warnings before use. Keep the carton. It contains important information.

Ranitidine Tablets USP, 75 mg - container label



Ranitidine Tablets USP, 75 mg - Container CartonLabel



RANITIDINE

ranitidine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55111-131
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ranitidine hydrochloride (UNII: BK76465IHM) (ranitidine - UNII:884KT10YB7)	ranitidine	75 mg

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSES (UNII: 3NXW29V3WO)	
magnesium stearate (UNII: 70097M6I30)	
cellulose, microcrystalline (UNII: OP1R32D61U)	
Polyethylene Glycol, Unspecified (UNII: 3WJQ0SDW1A)	
ferrosoferric oxide (UNII: XM0M87F357)	
titanium dioxide (UNII: 15FIX9V2JP)	

Product Characteristics

Color	PINK	**Score**	no score
Shape	ROUND	**Size**	7mm
Flavor		**Imprint Code**	P75
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55111-131-30	1 in 1 CARTON	03/01/2000	
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:55111-131-60	1 in 1 CARTON	03/01/2000	
2		60 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:55111-131-80	1 in 1 CARTON	03/01/2000	
3		80 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:55111-131-90	1 in 1 CARTON	03/01/2000	
4		90 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:55111-131-04	1 in 1 CARTON	03/01/2000	
5		120 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:55111-131-37	1 in 1 CARTON	03/01/2000	
6		160 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:55111-131-79	1 in 1 CARTON	03/01/2000	
7		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
8	NDC:55111-131-14	2 in 1 CARTON	03/01/2000	
8		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
9	NDC:55111-131-81	3 in 1 CARTON	03/01/2000	
9		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
10	NDC:55111-131-45	1 in 1 CARTON	03/01/2000	
10		45 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	ANDA075294	03/01/2000	

Labeler - Dr. Reddy's Laboratories Limited (650562841)