

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

Dr.Reddy's

NDC 55111-131-60

Regular Strength

Ranitidine

Tablets USP, 75 mg

ACID REDUCER

PREVENTS & RELIEVES
heartburn associated with acid
indigestion and sour stomach

60 Tablets (60 doses)

TAMPER EVIDENT: DO NOT USE IF FOIL SEAL UNDER CAP PRINTED WITH "SEALED FOR YOUR PROTECTION" IS BROKEN OR MISSING
IMPORTANT: This label does not contain full product information. See carton for complete information. Read the directions, consumer information leaflet and warnings before use. Retain carton and leaflet for reference.

Drug Facts

Active ingredient (in each tablet) Purpose

Ranitidine 75 mg (as ranitidine

hydrochloride USP, 84 mg) 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 See warnings on carton

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

■ do not use if carton or printed foil under 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

DISTRIBUTED BY: Dr. Reddy's Laboratories, Inc.

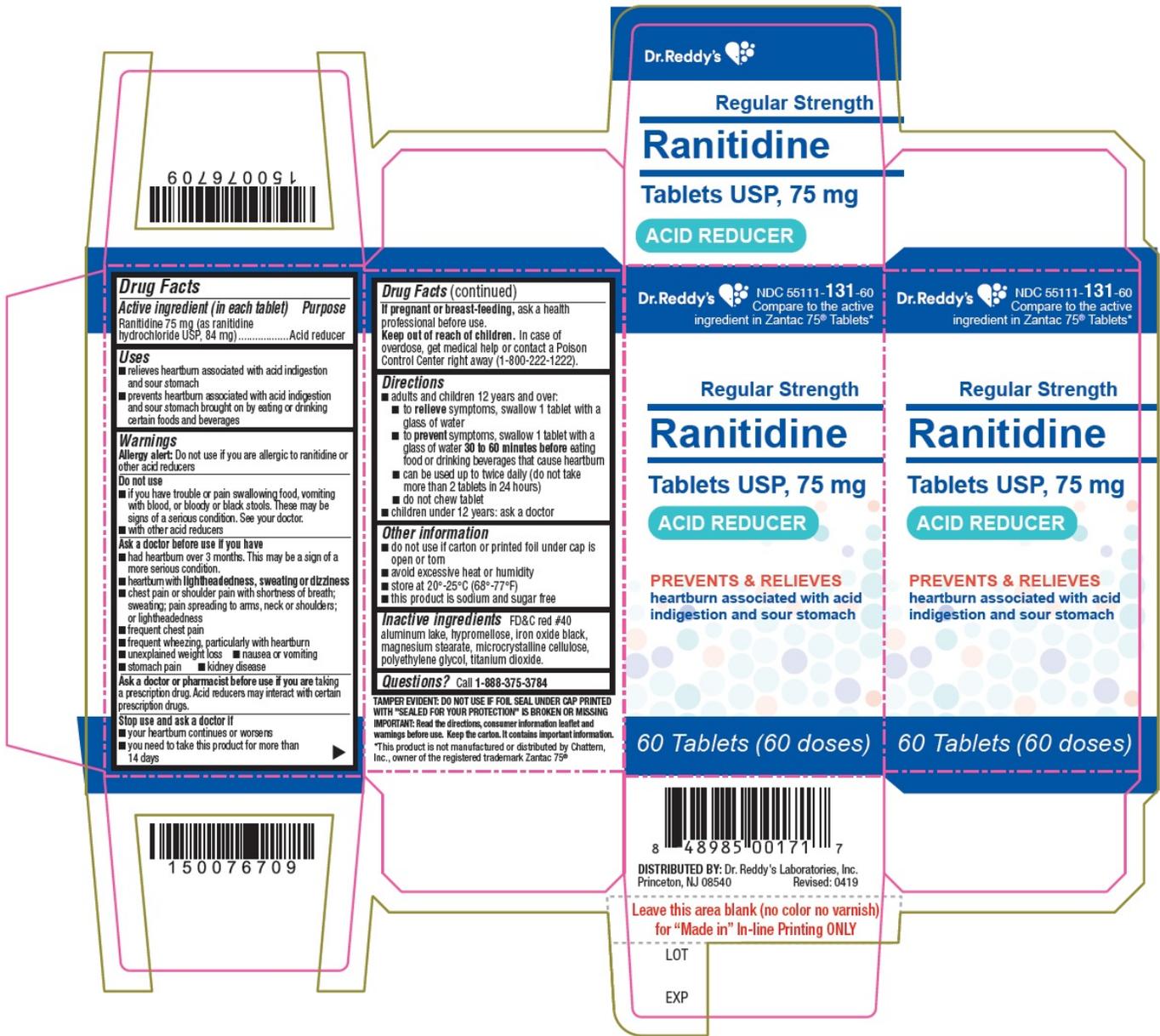
Princeton, NJ 08540

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LOT

EXP

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: 70097M6I3O)	
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		
	NDC:55111			

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)

Revised: 5/2025

Dr. Reddy's Laboratories Limited