RANITIDINE- ranitidine tablet, coated Dr. Reddy's Laboratories Inc.

Drug Facts

Active ingredient (in each tablet)

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

Purpose

Acid reducer

Use(s)

- 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
- if you have kidney disease, except under the advice and supervision of a doctor

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

Stop use and ask 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.

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

- do not use if printed foil under bottle cap is open or torn
- store at 20°-25°C (68°-77°F)
- avoid excessive heat or humidity
- protect from light
- this product is sodium and sugar free

Inactive ingredients

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

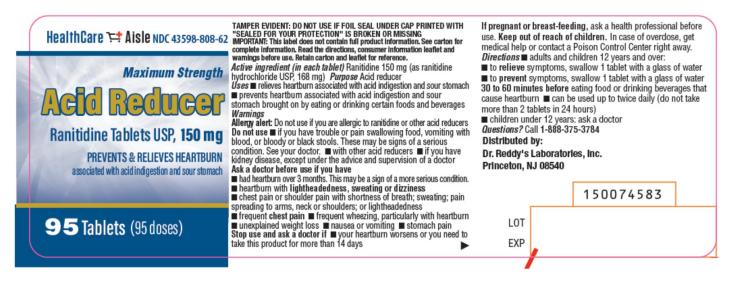
Questions?

Call 1-888-375-3784

Carton Label



Bottle Label



RANITIDINE

ranitidine tablet, coated

Product Information				
HUMAN OTC DRUG	Item Code (Source)	NDC:43598-808(NDC:55111-404)		
ORAL				

Active Ingredient/Active Moiety

Inare	edient	Name
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Basis of Strength Strength

Ranitidine Hydrochloride (UNII: BK76465IHM) (RANITIDINE - UNII:884KT10YB7)

RANITIDINE

150 mg

Inactive Ingr	redients
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mactive myredients			
Ingredient Name	Strength		
FD&C RED NO. 40 (UNII: WZB9127XOA)			
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)			
FERROSOFERRIC OXIDE (UNII: XM0M87F357)			
magnesium stearate (UNII: 70097M6I30)			
cellulose, microcrystalline (UNII: OP1R32D61U)			
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)			
titanium dioxide (UNII: 15FIX9V2JP)			

Product Characte	eristics
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Color	PINK	Score	no score
Shape	ROUND	Size	9mm
Flavor		Imprint Code	R150
Contains			

Packaging

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#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:43598-808- 62	1 in 1 CARTON	12/26/2018	09/30/2019		
1		95 in 1 BOTTLE; Type 0: Not a Combination Product				
2	NDC:43598-808- 65	1 in 1 CARTON	12/26/2018	09/30/2019		
2		220 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	ANDA078192	12/26/2018	

Labeler - Dr. Reddy's Laboratories Inc. (802315887)

Dr. Reddy's Laboratories Inc. Revised: 9/2019