

RANITIDINE 75- ranitidine tablets usp, 75mg tablet
The Kroger Co.

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

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

If pregnant or breast-feeding, ask a health professional before use.

Keep Out of Reach of Children.

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)
- 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.

Question Section

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

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

Package Label Principal Display Panel

Ranitidine Tablets USP, 75 mg - container label

NDC
30142-131-30



**Regular Strength
Heartburn
Relief 75**
Ranitidine Tablets USP, 75 mg
ACID REDUCER
Prevents and Relieves
Heartburn Associated
with Acid Indigestion
& Sour Stomach

30 TABLETS
(30 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)
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 1-800-632-6900
DISTRIBUTED BY THE KROGER CO., CINCINNATI, OHIO 45202
QUALITY GUARANTEE
www.kroger.com **GLUTEN FREE**

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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
Warning 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 2 months. This may be a sign of a more serious condition.
■ heartburn with **lightheadedness, swelling or dizziness**
■ chest pain or shoulder pain with shortness of breath, sweating, pain spreading to arms, neck or shoulders or lightheadedness
■ frequent wheezing, particularly with heartburn
■ unexplained weight loss
■ nausea or vomiting
■ stomach pain
Stop use and ask a doctor if ■ your heartburn continues or worsens
■ you need to take this product for more than 14 days
■ 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. (1-800-234-3229)

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
■ do not use if carton or printed foil under cap is open or torn

Drug Facts (continued)
■ avoid excessive heat or humidity
■ store at 20°-25°C (68°-77°F)
■ 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 1-800-632-6900

TAMPER EVIDENT: DO NOT USE IF FOIL SEAL UNDER CAP PRINTED WITH "SEALED FOR YOUR PROTECTION" IS BROKEN OR MISSING
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*Zantac 75 is a registered trademark of Boehringer Ingelheim Pharmaceuticals, Inc., Ridgebury, CT 06877. Boehringer Ingelheim Pharmaceuticals, Inc. is not affiliated with The Kroger Co. or this product.

GLUTEN FREE

COMPARE TO the active ingredient of ZANTAC 75 TABLETS
*See side panel

NDC 30142-131-30

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30 TABLETS (30 DOSES)

100% SIZE
12-digit UPCA
(non-suppressed)
For Position Only
0 11110 38507 9

DISTRIBUTED BY THE KROGER CO.
CINCINNATI, OHIO 45202
QUALITY GUARANTEE
www.kroger.com

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Leave this area blank (no color or warning) for "Made in" 8-line Printing ONLY

RANITIDINE 75

ranitidine tablets usp, 75mg tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:30142-131(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:30142-131-30	1 in 1 CARTON	03/31/2011	09/30/2019
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:30142-131-60	1 in 1 CARTON	03/31/2011	09/30/2019
2		60 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	03/31/2011	

Labeler - The Kroger Co. (006999528)

Revised: 9/2019

The Kroger Co.