RANITIDINE 150- ranitidine hydrochloride tablets 150mg tablet, coated
Walmart Stores Inc.

Drug Facts

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 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 diziness**
- 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 **1-888-287-1915**

TAMPER EVIDENT: DO NOT USE IF FOIL SEAL UNDER CAP PRINTED WITH "SEALED FOR YOUR PROTECTION" IS BROKEN OR MISSING

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

Ranitidine Tablets, 150 mg 24 count Carton

NDC 49035-404-34

equate™

Compare to Maximum Strength Zantac 150® Tablets Active Ingredient*

Maximum Strength

Ranitidine
Tablets, 150 mg

Acid Reducer

- PREVENTS AND RELIEVES HEARTBURN associated with acid indigestion and sour stomach

150 mg 24 TABLETS (24 DOSES)

Ranitidine Tablets, 150 mg 24 count Bottle

equate™ NDC 49035-404-34

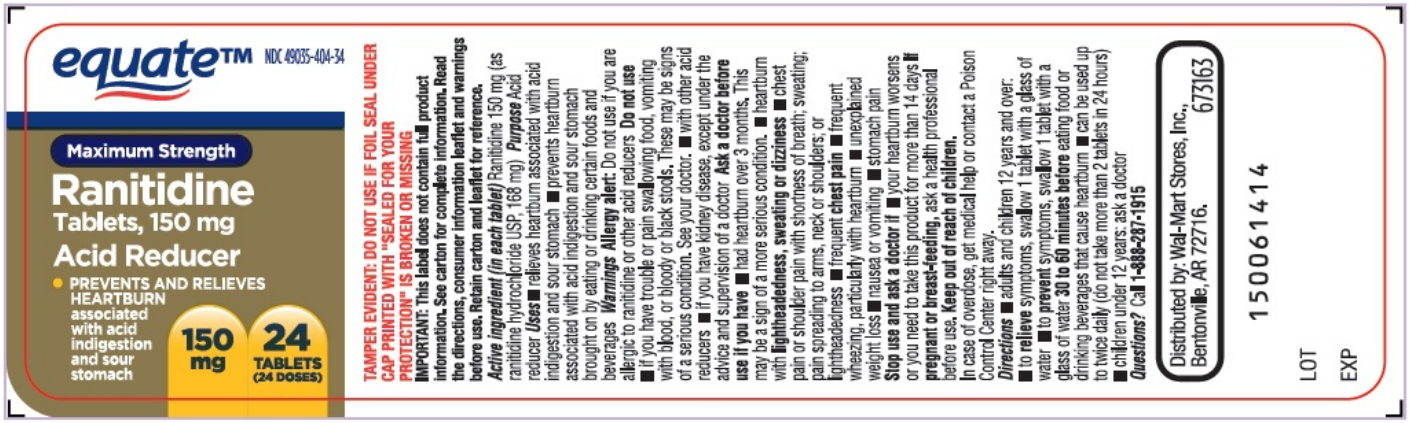
Maximum Strength

Ranitidine
Tablets, 150 mg

Acid Reducer

- PREVENTS AND RELIEVES HEARTBURN associated with acid indigestion and sour stomach

150 mg 24 TABLETS (24 DOSES)



RANITIDINE 150

ranitidine hydrochloride tablets 150mg tablet, coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-404(NDC:55111-404)
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**
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 Characteristics

Color	PINK	**Score**	no score
Shape	ROUND	**Size**	9mm
Flavor		**Imprint Code**	R150
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-404-34	1 in 1 CARTON	01/05/2010	09/30/2019
1		24 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:49035-404-61	1 in 1 CARTON	01/05/2010	09/30/2019
2		65 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:49035-404-13	2 in 1 CARTON	06/08/2018	09/30/2019
3		65 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:49035-404-65	1 in 1 CARTON	01/05/2010	09/30/2019
4		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	01/05/2010	

Labeler - Walmart Stores Inc. (051957769)

Revised: 9/2019 Walmart Stores Inc.