ACID REDUCER- ranitidine tablet

McKesson

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 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

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

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)
- children under 12 years: ask a doctor

OTHER INFORMATION

- TAMPER EVIDENT: DO NOT USE IF THE CARTON OR PRINTED FOIL UNDER CAP IS OPEN OR TORN.
- store at 20° 25° C (68° 77° F)
- avoid excessive heat or humidity
- this product is sodium and sugar free

INACTIVE INGREDIENTS

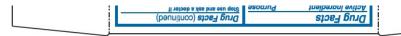
Colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide red, magnesium stearate, microcrystalline cellulose, polyethylene glycol, talc, titanium dioxide

QUESTIONS?

call 1-800-406-7984

PRINCIPAL DISPLAY PANEL sunmark[®] *COMPARE TO ZANTAC 150[®] ACTIVE INGREDIENT NDC 49348-026-04 Ranitidine Tablets, USP 150 mg acid reducer Prevents and Relieves Heartburn Associated with Acid Indigestion & Sour Stomach Maximum Strength 24 TABLETS Distributed By McKesson 5104606 / 0813





ACID REDUCER	2								
ranitidine tablet	-								
Product Information	n								
Product T ype	HUMAN OTC DRUG Item Code (Source) NI				ND	DC:49348-026			
Route of Administration	n 0								
Active Ingredient/Active Moiety									
		redient Name			Basis of St	rength	Strength		
RANITIDINE HYDRO CHL	L ORIDE (UNII: E	K76465IHM) (RANITIE	DINE - UNI	II:884KT10YB7)	RANITIDINE		150 mg		
Inactive Ingredients									
	Ingredient Name						Strength		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)									
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)									
HYPROMELLOSES (UNII: 3NXW29V3WO)									
FERRIC OXIDE RED (UNII: 1K09F3G675)									
MAGNESIUM STEARATE (UNII: 70097M6I30)									
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)									
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)									
TALC (UNII: 7SEV7J4R1U)									
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)									
Product Characteris	stics								
Color p	pink Score			no scoi		2			
Shape C	OVAL (Oval Sha	VAL (Oval Shaped) Size		Size		12mm	12mm		
Flavor		Imprint Code		Imprint Code	9 R				
Contains									
Packaging									
# Item Code	Packa	ge Description	Mar	keting Start Date	Mark	eting Er	nd Date		
1 NDC:49348-026-04	1 in 1 CART	ON							
1	24 in 1 BOT								
2 NDC:49348-026-54	1 in 1 CART								
2	65 in 1 BOT	TLE							
Marketing Information									
Marketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End Date									

0 1/0 7/20 14

Labeler - McKesson (177667227)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment

Name	Address	ID/FEI	Business Operations
Shasun Pharmaceuticals Limited		915786829	manufacture(49348-026)

Revised: 1/2014

McKesson