BERKLEY AND JENSEN HEARTBURN TREATMENT- esomeprazole capsule, delayed release

BJWC

BJWC Heartburn Treatment Drug Facts

Active ingredient (in each capsule)

Esomeprazole 20 mg

(Each delayed-release capsule corresponds to 22 mg esomeprazole magnesium dihydrate)

Purpose

Acid reducer

Uses

- treats frequent heartburn (occurs **2 or more** days a week)
- not intended for immediate relief of heartburn; this drug may take 1 to 4 days for full effect

Warnings

Allergy alert: Do not use if you are allergic to esomeprazole

Do not use if you have:

- trouble or pain swallowing food, vomiting with blood, or bloody or black stools
- 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**

These may be signs of a serious condition. See your doctor.

Ask a doctor before use if you have

- had heartburn over 3 months. This may be a sign of a more serious condition.
- frequent wheezing, particularly with heartburn
- unexplained weight loss
- nausea or vomiting
- stomach pain

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
- you need to take more than 1 course of treatment every 4 months
- you get diarrhea
- you develop a rash or joint pain

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 18 years of age and older
- this product is to be used once a day (every 24 hours), every day for 14 days
- may take 1 to 4 days for full effect

14-Day Course of Treatment

- swallow 1 capsule with a glass of water before eating in the morning
- take every day for 14 days
- do not take more than 1 capsule a day
- swallow whole. Do not crush or chew capsules.
- do not use for more than 14 days unless directed by your doctor

Repeated 14-Day Courses (if needed)

- you may repeat a 14-day course every 4 months
- do not take for more than 14 days or more often than every 4 months unless directed by a doctor
- children under 18 years of age: ask a doctor before use. Heartburn in children may sometimes be caused by a serious condition.

Other information

- read the directions and warnings before use
- keep the carton. It contains important information.
- store at 20-25°C (68-77°F)

Inactive ingredients

FD&C blue no. 1, FD&C red no. 3, ferric oxide, gelatin, glyceryl monostearate, hypromellose, magnesium stearate, meglumine, methacrylic acid and ethyl acrylate copolymer dispersion, polyethylene glycol, polysorbate 80, shellac, sodium lauryl sulfate, sugar spheres, talc, titanium dioxide, triethyl citrate

Questions or comments?

1-800-934-1204

Package/Label Principal Display Panel

Compare to the active ingredient in Nexium® 24 HR

SEE NEW WARNING INFORMATION

HEARTBURN TREATMENT

24 HOUR

ESOMEPRAZOLE MAGNESIUM DELAYED RELEASE CAPSULES 20 mg

ACID REDUCER

MAY TAKE 1 TO 4 DAYS FOR FULL EFFECT

TREATS FREQUENT HEARTBURN

42 CAPSULES

THREE 14-DAY COURSES OF TREATMENT

CAPSULES

ACTUAL SIZE

100% MONEY-BACK GUARANTEE



BERKLEY AND JENSEN HEARTBURN TREATMENT esomeprazole capsule, delayed release Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:68391-898

Active Ingredient/Active Mo	iety

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I		Ingredient Name		Basis of Strength	Strength
I	ESOMEPRAZOLE (UNII: N3PA65	559FT) (ESOMEPRAZOLE	- UNII:N3PA6559FT)	ESOMEPRAZOLE	20 mg

Inactive Ingredients		
Ingredient Name	Strength	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
FD&C RED NO. 3 (UNII: PN2ZH5LOQY)		
FERRIC OXIDE RED (UNII: 1K09F3G675)		
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)		
GLYCERYL MONOSTEARATE (UNII: 230 OU9 XXE4)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MEGLUMINE (UNII: 6 HG8 UB2MUY)		
POLYSORBATE 80 (UNII: 6OZP39ZG8H)		
SODIUM LAURYL SULFATE (UNII: 368GB5141J)		
TALC (UNII: 7SEV7J4R1U)		
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)		
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
SHELLAC (UNII: 46 N10 7B710)		

Product Characteristics			
Color	BLUE (opaque)	Score	no score
Shape	CAPSULE (oblong)	Size	14mm
Flavor		Imprint Code	L898
Contains			

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:68391-898-03	3 in 1 CARTON	09/25/2017		
1	14 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	ANDA207193	09/25/2017	

Revised: 1/2020 BJWC