## IBUTAB - ibuprofen tablet Zee Medical Inc

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### **Drug Facts**

Active Ingredient/Tablet Ibuprofen 200 mg (NSAID\*) \*nonsteroidal anti-inflammatory drug Purpose-Pain Reliever/Fever Reducer

#### Uses

- temporarily relieves minor aches and pains due to:
- headache the common cold muscular aches
- backache toothache minor pain of arthritis
- menstrual cramps
- temporarily reduces fever

#### Directions

- do not take more than directed
- the smallest effective dose should be used
- do not take longer than 10 days, unless directed by a doctor (see Warnings)
- adults: take 1 tablet every 4 to 6 hours while symptoms persist
- if pain or fever does not respond to 1 tablet, 2 tablets may be used
- do not exceed 6 tablets in 24 hours
- children under 12 years: ask a doctor

## Warnings

Allergy alert: Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin. Symptoms may include:

- hives facial swelling asthma (wheezing) shock
- rash skin reddening blisters

If an allergic reaction occurs, stop use and seek medical help right away.

Stomach bleeding warning: This product contains a nonsteroidal anti-inflammatory drug (NSAID), which may cause severe stomach bleeding. The chance is higher if you

- are age 60 or older
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
- have 3 or more alcoholic drinks every day while using this product
- take more or for a longer time than directed

#### Do not use

- if you have ever had an allergic reaction to any other pain reliever/fever reducer
- right before or after heart surgery

Ask a doctor before use if

- stomach bleeding warning applies to you
- you have a history of stomach problems, such as heartburn
- you have problems or serious side effects from taking pain relievers or fever reducers

- you have high blood pressure, heart disease, liver cirrhosis, kidney disease, or asthma
- you are taking a diuretic

Ask a doctor or pharmacist before use if you are

- under a doctor's care for any serious condition
- taking aspirin to prevent heart attack or stroke, because ibuprofen may decrease this benefit of aspirin
- taking any other drug

When using this product

- take with food or milk if stomach upset occurs
- the risk of heart attack or stroke may increase if you use more than directed or for longer than directed

Stop use and ask a doctor if

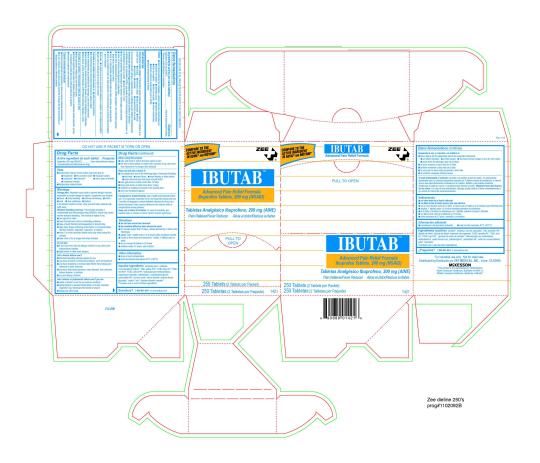
- you experience any of the following signs of stomach bleeding:
- feel faint vomit blood have bloody or black stools
- have stomach pain that does not get better
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present in the painful area
- any new symptoms appear

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use ibuprofen during the last 3 months of pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

KEEP OUT OF REACH OF CHILDREN. In case of overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Inactive ingredients carnauba wax\*, cellulose, croscarmellose sodium\*, DC yellow 10\*, FDC blue 2\*, FDC red 40\*, FDC yellow 6\*, hydroxypropyl methylcellulose, magnesium stearate, maltodextrin\*, polyethylene glycol\*, polysorbate 80\*, red iron oxide, silicon dioxide, sodium starch glycolate\*, starch\*, talc\*, titanium dioxide, triacetin\*
\*Contains one or more of these ingredients.

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# **IBUTAB**

ibuprofen tablet

Product Information			
Product Type	HUMAN OTC DRUG LABEL	Item Code (Source)	NDC:35418-750

Route of Administration ORAL DEA Schedule

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
IBUPRO FEN (IBUPRO FEN)	IBUPROFEN	200 mg	

Inactive Ingredients			
Ingredient Name	Strength		
CARNAUBA WAX			
CELLULO SE, MICRO CRYSTALLINE			
CROSCARMELLOSE SODIUM			
D&C YELLOW NO. 10			
FD&C BLUE NO. 2			
FD&C RED NO. 40			
FD&C YELLOW NO. 6			
MAGNESIUM STEARATE			

MALTO DEXTRIN	
POLYETHYLENE GLYCOL 300	
POLYSORBATE 80	
SILICON DIO XIDE	
SODIUM STARCH GLYCOLATE TYPE A POTATO	
STARCH, CORN	
TALC	
TITANIUM DIO XIDE	
TRIACETIN	

Product Characteristics			
Color	brown (chocolate brown)	Score	no score
Shape	ROUND (IBU;200)	Size	110 mm
Flavor		Imprint Code	IBU;200
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:35418-750-69	500 in 1 CARTON			
1	NDC:35418-750-67	125 in 1 CARTON			
1	NDC:35418-750-68	50 in 1 CARTON			
1	NDC:35418-750-02	2 in 1 PACKET			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA079129	06/12/2012	

# Labeler - Zee Medical Inc (009645623)

# Registrant - Ultra Seal Corporation (085752004)

Establishment				
Name	Address	ID/FEI	Business Operations	
Ultra Seal Corporation		085752004	repack(35418-750)	

Revised: 6/2012 Zee Medical Inc