

**IBUPROFEN- ibuprofen tablet, film coated**  
**Publix Super Markets Inc**

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**Publix Super Markets, Inc. Ibuprofen Caplets Drug Facts**

**Active ingredient (in each caplet)**

Ibuprofen 200 mg (NSAID)\*

\*nonsteroidal anti-inflammatory drug

**Purposes**

Pain reliever/fever reducer

**Uses**

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

**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
- skin reddening
- rash
- blisters

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

**Stomach bleeding warning:** This product contains an 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

**Heart attack and stroke warning:** NSAIDs, except aspirin, increase the risk of heart attack, heart failure, and stroke. These can be fatal. The risk is higher if you use more than directed or for longer than directed.

### **Do not use**

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

### **Ask a doctor before use if**

- you have problems or serious side effects from taking pain relievers or fever reducers
- the stomach bleeding warning applies to you
- you have a history of stomach problems, such as heartburn
- you have high blood pressure, heart disease, liver cirrhosis, kidney disease, asthma, or had a stroke
- you are taking a diuretic

### **Ask a doctor or pharmacist before use if you are**

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

### **When using this product**

- take with food or milk if stomach upset occurs

### **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
- you have symptoms of heart problems or stroke:
  - chest pain
  - trouble breathing
  - weakness in one part or side of body
  - slurred speech
  - leg swelling

- 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 at 20 weeks or later in 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 (1-800-222-1222).

**Directions**

- **do not take more than directed**
- **the smallest effective dose should be used**

adults and children 12 years and older	<ul style="list-style-type: none"> <li>• take 1 caplet every 4 to 6 hours while symptoms persist</li> <li>• if pain or fever does not respond to 1 caplet, 2 caplets may be used</li> <li>• do not exceed 6 caplets in 24 hours, unless directed by a doctor</li> </ul>
children under 12 years	<ul style="list-style-type: none"> <li>• ask a doctor</li> </ul>

**Other information**

- read all warnings and directions before use
- store between 68-77°F (20-25°C)
- avoid high humidity and excessive heat above 104°F (40°C)
- see end panel for lot number and expiration date

**Inactive ingredients**

colloidal silicon dioxide, corn starch, croscarmellose sodium, FD&C red no. 40 aluminum lake, FD&C yellow no. 6 aluminum lake, iron oxides, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, stearic acid, talc, titanium dioxide

**Principal Display Panel**

SEE REVISED WARNING

P

ibuprofen caplets\*\*

IBUPROFEN TABLETS, 200 mg

PAIN RELIEVER/FEVER REDUCER (NSAID)

ACTUAL SIZE

100 COATED CAPLETS\*\*

\*\*CAPSULE-SHAPED TABLETS

Compare to Motrin<sup>®</sup> IB active ingredient

DO NOT USE IF PAINED SEAL UNDER CAP  
ISBROKEN OR MISSING

Keep this box for important information before using.

**Drug Facts (continued)**

**Directions**  
 ■ do not take more than directed  
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 COMPLETE SATISFACTION OR YOUR MONEY BACK

**Publix.**



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Important: Read all product information before using.

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 \*nonsteroidal anti-inflammatory drug

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**P**

**ibuprofen caplets\*\***

IBUPROFEN TABLETS, 200 mg  
PAIN RELIEVER/FEVER REDUCER (NSAID)

**100** COATED CAPLETS\*\*  
\*\*CAPSULE-SHAPED TABLETS

Compare to Motrin® IB active ingredient\*\*\*



ACTUAL SIZE

NDC 56062-517-78  
SEE REVISED WARNING

51778 63 C6

<b>IBUPROFEN</b>			
ibuprofen tablet, film coated			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:56062-517
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	200 mg

**Inactive Ingredients**

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

**Product Characteristics**

Color	ORANGE	Score	no score
Shape	OVAL	Size	15mm
Flavor		Imprint Code	I2
Contains			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:56062-517-71	1 in 1 CARTON	05/04/2006	
1		50 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:56062-517-78	1 in 1 CARTON	05/03/2006	
2		100 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	ANDA077349	05/03/2006	

