

**IBUPROFEN- ibuprofen tablet, film coated**  
**Better Living Brands, LLC**

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**Signature Care 44-393**

***Active ingredient (in each orange caplet)***

Ibuprofen USP,  
200 mg (NSAID)\*  
\*nonsteroidal anti-inflammatory drug

***Purpose***

Pain reliever/fever reducer

***Uses***

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

- take a blood thinning (anticoagulant) or steroid drug
- take more or for a longer time than directed
- 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
- are age 60 or older
- have had stomach ulcers or bleeding problems

**Do not use**

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

**Ask a doctor before use if**

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

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

- under a doctor's care for any serious condition
- taking aspirin for 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:
  - vomit blood
  - feel faint
  - 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 complication during delivery.

**Keep out of reach of children**

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

***Directions***

- **do not take more than directed**
- **the smallest effective dose should be used**
- adults and children 12 years and over: take 1 caplet every 4 to 6 hours while symptoms persist
- if pain or fever does not respond to 1 caplet, 2 caplets may be used
- do not exceed 6 caplets in 24 hours, unless directed by a doctor
- children under 12 years: ask a doctor

***Other information***

- store between 20°-25°C (68°-77°F)
- avoid excess heat 40°C (104°F)
- use by expiration date on package

***Inactive ingredients***

carnauba wax, cellulose, corn starch, FD&C yellow #6 aluminum lake, fumed silica gel, hypromellose, lactose, magnesium stearate, polydextrose, polyethylene glycol, sodium starch glycolate, stearic acid, titanium dioxide

***Questions or comments?***

**Call 1-800-426-9391** 8:30 AM-4:00 PM ET, Monday-Friday

***Principal Display Panel***

***Signature™***

care  
Quality Guaranteed

**COMPARE TO** Motrin® IB Caplets active ingredient†

NDC 21130-393-14

**Ibuprofen**

IBUPROFEN TABLETS USP, 200 mg

Pain Reliever / Fever Reducer

**(NSAID)**

Actual Size

**500 COATED  
CAPLETS**

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS  
BROKEN OR MISSING**

†This product is not manufactured or distributed by McNeil Consumer Healthcare, distributors of Motrin® IB Caplets.

50844 REV0616A39314

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P.O. BOX 99, PLEASANTON, CA 94566-0009


1-888-723-3929 [www.betterlivingbrandsLLC.com](http://www.betterlivingbrandsLLC.com)

OUR PROMISE, QUALITY & SATISFACTION

100% GUARANTEED OR YOUR MONEY BACK

RD 16277

**COMPARE TO**  
Motrin® IB Caplets  
active ingredient†  
NDC 21130-393-14



Quality  
Guaranteed

**Drug Facts** TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

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\*nonsteroidal anti-inflammatory drug

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     ■ backache      ■ toothache  
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**Drug Facts (continued)**

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     ■ feel faint   ■ have bloody or black stools  
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**Drug Facts (continued)**

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**Ibuprofen**


IBUPROFEN TABLETS USP, 200 mg  
Pain Reliever/Fever Reducer  
(NSAID)

Actual Size



500 COATED CAPLETS

RD 16277



S1714

21130-393-14

100% GUARANTEED OR YOUR MONEY BACK

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 1-888-273-3939 www.betterlivingbrandsllc.com  
 OUR PROMISE: QUALITY & SATISFACTION

Signature Care 44-393

IBUPROFEN			
ibuprofen tablet, film coated			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-393
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
IBUPROFEN (UNII: WK2XYH10QM) (IBUPROFEN - UNII:WK2XYH10QM)	IBUPROFEN	200 mg	
Inactive Ingredients			
Ingredient Name	Strength		
CARNAUBA WAX (UNII: R12CBM0EIZ)			
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)			
HYPROMELLOSES (UNII: 3NXW29V3WO)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
POLYDEXTROSE (UNII: VH2XOU12IE)			
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)			
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)			
STEARIC ACID (UNII: 4ELV7Z65AP)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
Product Characteristics			
Color	ORANGE	Score	no score
Shape	OVAL	Size	14mm

<b>Flavor</b>		<b>Imprint Code</b>	44;393	
<b>Contains</b>				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-393-12	1 in 1 CARTON	03/01/1999	
1		100 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:21130-393-15	1 in 1 CARTON	03/01/1999	
2		50 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:21130-393-14	500 in 1 BOTTLE; Type 0: Not a Combination Product	03/01/1999	
<b>Marketing Information</b>				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA075139	03/01/1999		

**Labeler** - Better Living Brands, LLC (009137209)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	PACK(21130-393)

### Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	MANUFACTURE(21130-393)

Revised: 1/2017

Better Living Brands, LLC