IBUPROFEN- ibuprofen tablet, film coated CVS Pharmacy

CVS 44-392

Active ingredient (in each orange tablet)

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

Purpose

Pain reliever/fever reducer

Uses

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

Warnings

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

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

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

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 any other pain reliever/fever reducer
- right before or after heart surgery

Ask a doctor before use if

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

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
 - leg swelling
 - slurred speech
 - weakness in one part or side of body
- 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 (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 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, unless directed by a doctor
- children under 12 years: ask a doctor

Other information

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

Inactive ingredients

carnauba wax, colloidal silicon dioxide, corn starch, FD&C yellow #6 aluminum lake, hypromellose, lactose anhydrous, magnesium stearate, microcrystalline cellulose, 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

♥ CVS Health™

Compare to the active ingredient in Motrin[®] IB[†]

Coated Tablets

NDC 59779-392-14

Ibuprofen IBUPROFEN TABLETS USP, 200 mg

Pain reliever, Fever reducer **(NSAID)**

Actual Size

500 COATED TABLETS

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

[†]This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Motrin[®] IB.

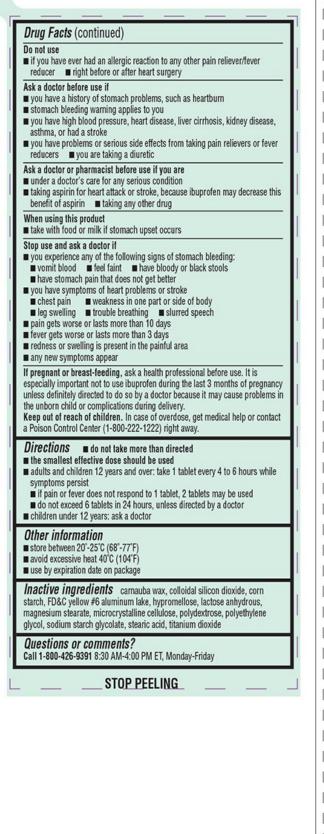
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CVS 44-392

ibuprofen tablet, film coated

Product Inform	ation						
Product T ype		HUMAN OTC DRUG	C DRUG Item Code (Source)		NDC:59779-392		
Route of Administr							
Active Ingredie	nt/Active Moi	ety					
Ingredient Name					Basis of Strength		
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)					IBUPRO FEN		200 mg
Inactive Ingredi	ents						
		Ingredient	Name				Strength
CARNAUBA WAX (U	NII: R12CBM0EIZ)					
STARCH, CORN (UNII: 08232NY3SJ)							
FD&C YELLOW NO.6 (UNII: H77VEI93A8)							
MAGNESIUM STEAI		,					
POLYDEXTROSE (U							
STEARIC ACID (UNI							
TITANIUM DIO XIDE							
HYPROMELLOSE, U							
ANHYDRO US LACT							
		CIFIED (UNII: 3WJQ0S	DW1A)				
SILICON DIOXIDE (
		E (UNII: OP1R32D61U)					
SODIUM STARCH G	LICOLATEIII	PE A POTATO (UNII:	5850J3G2A2)				
Product Charac							
Color	ORANO		Score		no score		
Shape	ROUND Size			10 mm			
Flavor Imprint Code 44;392							
Contains							
Packaging		Package Description			rketing Start	Mar	keting End
		Package Descrip	tion	Ma	Date		Date
# Item Code	1 in 1 CARTON	Package Descrip	tion	Ma 03/01	Date	02/20/2	
 # Item Code 1 NDC:59779-392- 08 1 		Package Descrip		03/01	Date	02/20/2	
 # Item Code 1 NDC:59779-392- 08 1 NDC:50770-202 	24 in 1 BOTTLE, Product 1 in 1 CARTON	PLASTIC; Type 0: No	t a Combinatio	0 3/0 1 on 03/0 1	Date /1999	02/20/2 10/20/2	0 19
 # Item Code 1 NDC:59779-392- 08 1 NDC:59779-392- 15 2 NDC:59779-392- 15 	24 in 1 BOTTLE, Product 1 in 1 CARTON		t a Combinatio	0 3/0 1 on 03/0 1	Date /1999		0 19
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4		500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	0 3/0 1/19 9 9				
Marketing Information							
N	Aarketing Categor	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
	NDA	ANDA075139	03/01/1999				
A	NDA						

Labeler - CVS Pharmacy (062312574)

Name	Addres	s	ID/FEI	Business Operations	
NK International, Inc.		0381	54464	PACK(59779-392)	
Establishment					
Name	Address	ID/	FEI	Business Operations	
LNK International, Inc.		83286789	4 MAN	UFACTURE(59779-392)	
Establishment					
Name	Addres	s	ID/FEI	Business Operations	
NK International, Inc.		96762	6305	PACK(59779-392)	

Lstablishillent						
	Name	Address	ID/FEI	Business Operations		
	LNK International, Inc.		832867837	PACK(59779-392)		

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	PACK(59779-392)

Revised: 4/2020

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