# UP AND UP IBUPROFEN- ibuprofen tablet, film coated Target Corporation

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## Target Corporation Ibuprofen Tablets, 200 mg 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 chances are 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 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 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 (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:

- 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

- read all warnings and directions before use
- store at 20-25°C (68-77°F)
- avoid high humidity and excessive heat above 40°C (104°F)

## **Inactive ingredients**

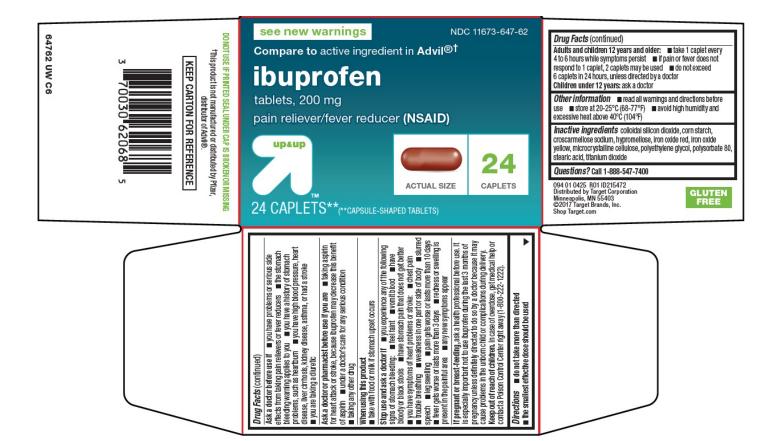
colloidal silicon dioxide, corn starch, croscarmellose sodium, hypromellose, iron oxide red, iron oxide yellow, microcrystalline cellulose, polyethylene glycol, polysorbate 80, stearic acid, titanium dioxide

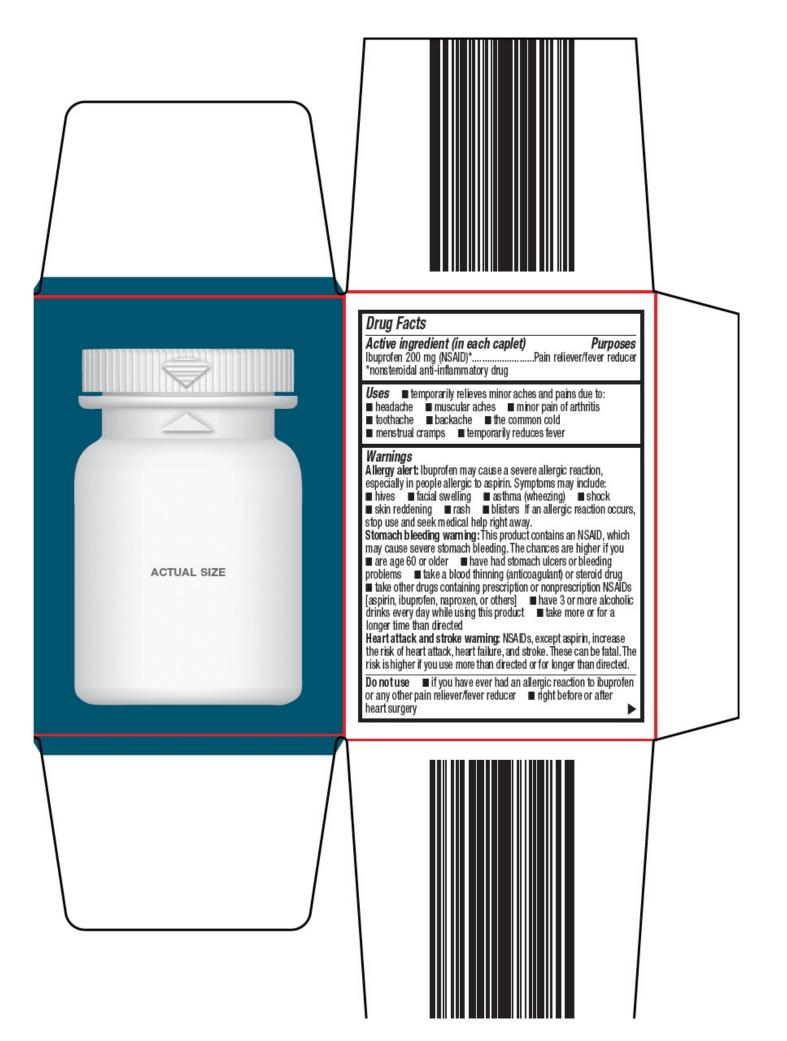
#### Questions?

Call 1-888-547-7400

#### **Principal Display Panel**

see new warnings Compare to active ingredient in Advil® ibuprofen tablets, 200 mg pain reliever/fever reducer (NSAID) ACTUAL SIZE 24 CAPLETS 24 CAPLETS\*\* (\*\*CAPSULE-SHAPED TABLETS)





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ibι	ıprofen tablet, film	n coated										
P	roduct Informa	tion										
D	roduct Type		ΗΙΜΔΝ	OTC DRU	IG	Itom (	oda (Sau	rca)	NDC:116	73-647		
Product Type						ite in C	tem Code (Source)		100.11075-047			
Route of Administration			ORAL									
-		<i></i>										
A	ctive Ingredien											
Ingredient Name								Basis of St	rength	Strength		
IE	BUPROFEN (UNII: WI	K2XYI10Q	M) (IBUPROFEN	I - UNII:WK	K2XYI10QM)			IBUPROFEN		200 mg		
т	·····											
11	nactive Ingredie	ents										
			-	gredient	Name					Strength		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)												
	FARCH, CORN (UNI			Π140)								
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48) HYPROMELLOSES (UNII: 3NXW29V3WO)												
	ERRIC O XIDE RED (											
				)								
	FERRIC OXIDE YELLOW (UNII: EX43802MRT)   MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)											
	POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)											
POLYSORBATE 80 (UNII: 60ZP39ZG8H)												
S	STEARIC ACID (UNII: 4ELV7Z65AP)											
T	ITANIUM DIO XIDE (	(UNII: 15FI	X9V2JP)									
Product Characteristics												
С	olor	]	BROWN		Score							
Shape			OVAL	L Si		Size			15mm			
Flavor			Imprint Code			12						
Contains												
P	ackaging											
#	# Item Code		Package Description				Marketi	ng Start Date	Market	ing End Date		
1	NDC:11673-647-62	1 in 1 CARTON			05/28/2009			9				
1		24 in 1 BOTTLE; Type 0: Not a Com										
	NDC:11673-647-85	1 in 1 CA					06/12/200	9				
2			BOTTLE; Type 0									
	NDC:11673-647-90		BOTTLE; Type (	): Not a Co	ombination Pro	duct	06/12/200		04/09/20	15		
	NDC:11673-647-82	1 in 1 CA			and the state of the the	J.,	02/23/201	7				
4		200 in 11	BOTTLE; Type (	): Not a Co	ombination Pro	duct						

Marketing Information									
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date						
ANDA	ANDA072096	05/28/2009							

Labeler - Target Corporation (006961700)

Revised: 12/2019

Target Corporation