

IBUPROFEN- ibuprofen capsule, liquid filled

H E B

Drug Facts

Active ingredient (in each capsule)

Solubilized ibuprofen equal to 200 mg ibuprofen (NSAID)*

(present as the free acid and potassium salt)

*nonsteroidal anti-inflammatory drug

Purpose

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
 - muscular 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 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 problems or serious side effects from taking pain relievers or fever reducers
- 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

- 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 problem or stroke:
 - chest pain
 - slurred speech
 - trouble breathing
 - leg swelling
 - 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 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 (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 capsule every 4 to 6 hours while symptoms persist
 - if pain or fever does not respond to 1 capsule, 2 capsules may be used
 - do not exceed 6 capsules in 24 hours, unless directed by a doctor
- children under 12 years: ask a doctor

Other information

- **each capsule contains:** potassium 20 mg
- read all warnings and directions before use keep carton.
- store at 20° to 25°C (68° to 77°F)
- avoid excessive heat above 40°C (104°F). Protect from light.
- swallow whole; do not crush, chew, or dissolve

Inactive ingredients

FD&C blue #1, gelatin, pharmaceutical ink, polyethylene glycol, potassium hydroxide, purified water, sorbitan, sorbitol

Questions & Comments

Call toll free: **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal Display Panel

Compare to Advil® Liquid-Gels® active ingredient**

Ibuprofen

Capsules, 200 mg

Pain Reliever/Fever Reducer (NSAID)

For Body Aches and Pain

CAPSULES

(†Liquid Filled Capsules)

****This product is not manufactured or distributed by GlaxoSmithKline Consumer Healthcare Holdings (US) LLC, distributor of Advil® Liqui-Gels®.**

MADE WITH PRIDE AND CARE FOR H-E-B®, SAN ANTONIO, TX 78204

Product Label

Compare to Advil® Liqui-Gels® active ingredient*

NDC 37808-699-30

H-E-B

Ibuprofen
Capsules, 200 mg

Pain Reliever/Fever Reducer (NSAID)

For Body Aches and Pain

actual size

300 CAPSULES
SOFTGELS† ('Liquid Filled Capsules')

IBUPROFEN 200mg SOFTGELS

Drug Facts

Active ingredient (in each capsule)
Sulfonized ibuprofen equal to 200 mg ibuprofen (NSAID)*. Pain reliever/fever reducer (present as the free acid and potassium salt). Nonsteroidal anti-inflammatory drug.

Uses

- temporarily relieves minor aches and pains due to:
 - headache
 - minor pain of arthritis
 - toothache
 - 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)
- skin rash
- dizziness
- stomach bleeding

If an allergic reaction occurs, stop use and seek medical help right away. If you have had stomach bleeding, the chance is higher if you:

- are age 60 or older
- have had stomach bleeding
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
- have a blood thinning (anticoagulant or stomach or bleeding) problem
- take more than 3 or more alcoholic drinks every day while using this product
- take more than 1 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

Drug Facts (continued under label)

*This product is not manufactured or distributed by BayerSmithKline Consumer Healthcare Holdings (US) LLC, distributor of Advil® Liqui-Gels®.

MADE WITH PRIDE AND CARE FOR H-E-B® SAN ANTONIO, TX 78264

100% GUARANTEE
promise

If you aren't completely satisfied with this product, we'll be happy to replace it or refund your money. You have our word on it.

FD-036 1000266 Rev 02/02
41071-2/06

Exp. Date: 0 4 1220 28430 6

LOT: 01222

PERFECT PRICE

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

Drug Facts (continued)

Ask a doctor before use if

- the stomach bleeding warning applies to you
- you have problems or serious side effects from taking pain relievers or fever reducers
- 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

- 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, like 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
 - slurred speech
 - trouble breathing
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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 (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:
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Other information

- each capsule contains: potassium 20 mg
- all warnings are directions before use
- store at 20° to 25°C (68° to 77°F)
- avoid excessive heat above 40°C (104°F). Protect from light.
- swallow whole; do not crush, chew, or dissolve

Inactive ingredients FDAC blue #1, gelatin, pharmaceutical ink, polyethylene glycol, potassium hydroxide, purified water, sorbitol, xanthan

Questions or comments?

Call toll free: 1-877-755-8835 Monday-Friday 9AM-5PM EST

IBU PROFEN

ibuprofen capsule, liquid filled

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:37808-699
Route of Administration		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
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
GELATIN (UNII: 2G86QN327L)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)				
WATER (UNII: 059QF0KO0R)				
SORBITOL (UNII: 506T60A25R)				
SORBITAN (UNII: 6O92ICV9RU)				
Product Characteristics				
Color		blue (green)	Score	no score
Shape		CAPSULE	Size	18mm
Flavor			Imprint Code	IB200
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-699-80	1 in 1 BOX	03/31/2018	
1		80 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:37808-699-30	300 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/31/2018	
3	NDC:37808-699-20	1 in 1 BOX	03/31/2018	
3		20 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:37808-699-40	1 in 1 BOX	03/31/2018	
4		40 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

ANDA	ANDA078682	03/31/2018	
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Labeler - H E B (007924756)

Revised: 10/2023

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