

IBUPROFEN- ibuprofen capsule, liquid filled
Publix Super Markets Inc

Publix Super Markets, Inc. Ibuprofen Liquid Gels 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
- toothache
- backache
- menstrual cramps
- the common cold
- 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:

- 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 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 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 68-77°F (20-25°C)
- avoid excessive heat above 104°F (40°C)

Inactive ingredients

ammonium hydroxide, caprylic and capric acid triglycerides, FD&C green No. 3, gelatin, isopropyl alcohol, lecithin, macrogol/PEG 400, polyethylene glycol, polyvinyl acetate phthalate, potassium hydroxide, propylene glycol, purified water, SDA 35A alcohol, sorbitol sorbitan solution, titanium dioxide

Principal Display Panel

ibuprofen liquid gels**

SOLUBILIZED IBUPROFEN CAPSULES, 200 mg


PAIN RELIEVER/FEVER REDUCER (NSAID)

ACTUAL SIZE

80 LIQUID GELS**


****LIQUID-FILLED CAPSULES**


Compare to Advil® Liqui-Gels® active ingredient



ibuprofenliiquidgels^{**}


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





80 LIQUID GELS^{**}

****LIQUID-FILLED CAPSULES**



ACTUAL SIZE

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

NDC 56062-131-27

READ AND KEEP CARTON FOR COMPLETE WARNINGS AND INFORMATION

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temporarily relieves minor aches and pains due to:

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- menstrual cramps
- minor pain of arthritis
- temporarily reduces fever

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Drug Facts (continued)

Heart attack and stroke warning: NSAIDs, except aspirin, increase the risk of heart attack, heart failure, and stroke. These can be fatal. The risks are 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
- you have a history of stomach problems, such as heartburn
- you have high blood pressure, heart disease, liver disease, kidney disease, asthma, or had a stroke
- you are taking a diuretic

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 MADE IN CHINA
 PUBLIC SUPER MARKETS, INC.
 COMPLETE SATISFACTION OR YOUR MONEY BACK

Drug Facts (continued)

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DO NOT USE IF PRINTED SEAL UNDER CAP IS BROKEN OR MISSING



0 41415 41073 2

13127 63 C4

IBUPROFEN

ibuprofen capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:56062-131
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
AMMONIA (UNII: 5138Q19F1X)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ACETATE PHTHALATE (UNII: 58QVG85GW3)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SORBITOL (UNII: 506T60A25R)	
SORBITAN (UNII: 6O92ICV9RU)	

Product Characteristics

Color	GREEN (Light Green)	Score	no score
Shape	OVAL	Size	19mm
Flavor		Imprint Code	131
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:56062-131-58	1 in 1 CARTON	08/02/2017	
1		40 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:56062-131-27	1 in 1 CARTON	08/07/2017	
2		80 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:56062-131-06	1 in 1 CARTON	01/17/2018	
3		160 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	ANDA203599	08/02/2017	

Labeler - Publix Super Markets Inc (006922009)

Revised: 7/2022

Publix Super Markets Inc