

UP AND UP IBUPROFEN- ibuprofen capsule, liquid filled **Target Corporation**

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

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, sorbitol sorbitan solution, titanium dioxide

Questions?

Call 1-888-547-7400

Package/Label Principal Display Panel

see new warnings

Compare to active ingredient in Advil® Liquid-Gels®

ibuprofen liquid gelst

solubilized ibuprofen capsules, 200 mg

pain reliever/fever reducer (NSAID)

ACTUAL SIZE | 160 LIQUID GELS†

160 LIQUID GELS† (†LIQUID FILLED CAPSULES)

see new warnings

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

ibuprofen liquid gels[†]

solubilized ibuprofen capsules, 200 mg
pain reliever/fever reducer (NSAID)



ibuprofen liquid gels[†]

solubilized ibuprofen capsules, 200 mg
pain reliever/fever reducer (NSAID)



160 LIQUID GELS[†] (LIQUID-FILLED CAPSULES)

13108 LW 01

CODE AREA



DO NOT USE IF PRINTED OR UNDOCTORED FOR INFORMATION ONLY
Distributed by Parke-Davis Corporation
Warrenville, IL 60593
Made in Canada by
TMS Canada International Inc.

04101 0383 000
0-000004-01-010
3 770050 27540 3



READ AND KEEP CARTON FOR COMPLETE WARNING AND INFORMATION

Drug Facts

Active ingredient (in each capsule)
Solubilized ibuprofen liquid gel
200 mg ibuprofen (NSAID)
Pain reliever/fever reducer (NSAID)
(present as the free acid and potassium salt)
*as active ingredient in Advil® Liqui-Gels®

Purpose
Pain reliever
Fever reducer

Uses
It is primarily used to relieve pain and reduce fever 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 redness ● rash ● difficulty breathing
If you have any of these symptoms, 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 risks are higher if you take 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 or right before or after heart surgery

Ask a doctor before use if
● 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

Ask a doctor or pharmacist before use if you are
● under a doctor's care for any medical condition
● taking aspirin for heart attack or stroke, because ibuprofen may decrease the benefit of aspirin
● taking any other drug

Drug Facts (continued)

When using this product
● take with food or milk if stomach upsets occur
Stop use and ask a doctor if
● you experience any of the following signs of stomach bleeding:
● food taste ● vomit ● blood ● stools ● have bloody or black stools
● you have stomach pain that does not go away
● 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 ● numbness or tingling ● lasts more than 10 days
● fever ● gets worse or lasts more than 3 days
● asthma 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 directed 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 to 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
● not all heartburn and indigestion before use. Keep carbs.
● store at 20-25°C (68-77°F)
● avoid excessive heat above 40°C (104°F)

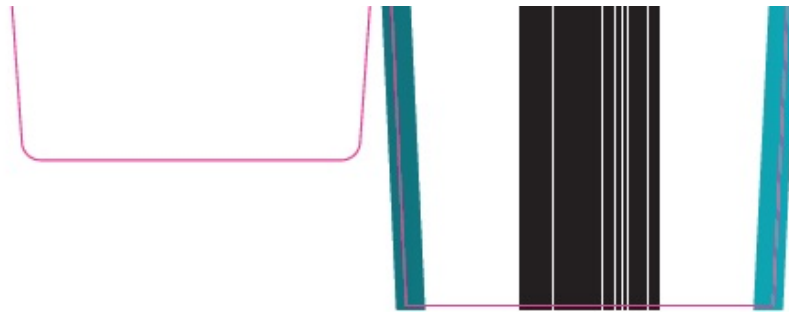
Inactive ingredients

ammonium hydroxide, caprylic acid, capric acid, hydroxyethylcellulose, FD&C green No. 3, gelatin, hydroxyethylcellulose, lactose, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, potassium hydroxide, propylene glycol, purified water, sodium hydroxide, sodium lauryl sulfate, titanium dioxide

Questions? Call 1-888-547-7400

**The products do not contain ibuprofen or are distributed by Parke-Davis, distributor of Advil® Liqui-Gels®.





UP AND UP IBUPROFEN

ibuprofen capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-514
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XY110QM) (IBUPROFEN - UNII:WK2XY110QM)	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

Marketing Start Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-514-06	1 in 1 CARTON	12/28/2018	02/28/2021
1		160 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:11673-514-58	1 in 1 CARTON	12/28/2018	07/31/2023
2		40 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:11673-514-76	1 in 1 CARTON	12/28/2018	12/28/2018
3		120 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:11673-514-27	1 in 1 CARTON	12/28/2018	
4		180 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:11673-514-87	300 in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2019	02/28/2021

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA203599	12/28/2018	

Labeler - Target Corporation (006961700)

Revised: 7/2022

Target Corporation