DYE FREE IBUPROFEN- ibuprofen tablet, film coated Walgreen Company

Walgreens 44-438-ADR

Active ingredient (in each white tablet)

Ibuprofen USP, 200 mg (NSAID)*
*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:

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

- take more or for a longer time than directed
- take a blood thinning (anticoagulant) or steroid drug
- are age 60 or older
- have had stomach ulcers or bleeding problems
- have 3 or more alcoholic drinks every day while using this product

take other drugs containing prescription or nonprescription NSAIDs [aspirin, ibuprofen, naproxen, or others]

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 a history of stomach problems, such as heartburn
- you are taking a diuretic
- 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

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
 - have bloody or black stools
 - vomit blood
 - have stomach pain that does not get better
- you have symptoms of heart problems or stroke:
 - chest pain
 - slurred speech
 - leg swelling
 - trouble breathing
 - 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 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)
- see end flap for expiration date and lot number

Inactive ingredients

colloidal silicon dioxide, corn starch, hypromellose, lactose anhydrous, magnesium stearate, microcrystalline cellulose, polydextrose, polyethylene glycol, povidone, sodium starch glycolate, stearic acid, triacetin

Questions or comments?

Call 1-800-426-9391 8:30 AM-4:00 PM ET, Monday-Friday

Principal Display Panel

Walgreens

WALGREENS PHARMACIST RECOMMENDED[†]

Compare to the active ingredient in Advil® Tablets††

NDC 0363-0438-99

Dye-Free

Ibuprofen

IBUPROFEN TABLETS USP, 200 MG / PAIN RELIEVER / FEVER REDUCER (NSAID)

100

TABLETS

ACTUAL SIZE

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

DISTRIBUTED BY: WALGREEN CO. 200 WILMOT RD., DEERFIELD, IL 60015 100% SATISFACTION GUARANTEED walgreens.com © 2021 Walgreen Co.

†Our pharmacists recommend the Walgreens brand. We invite you to compare to national brands. ††This product is not manufactured or distributed by PF Consumer Healthcare 1 LLC, owner of the registered trademark Advil® Tablets.

50844 ORG122143812



DYE FREE IBUPROFEN

ibuprofen tablet, film coated

| Product Information | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0363-0438 |
| Route of Administration | ORAL | | |
| | | | |
| | | | |

| Active Ingredient/Active Moiety | | | | |
|--|--------------------------|----------|--|--|
| Ingredient Name | Basis of Strength | Strength | | |
| IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM) | IBUPROFEN | 200 mg | | |

| STARCH, CORN (UNII: 08232NY3SJ) HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK) MAGNESIUM STEARATE (UNII: 70097M6I30) MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) POLYDEXTROSE (UNII: VH2XOU12IE) POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E) SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2) | Inactive Ingredients | |
|--|--|----------|
| STARCH, CORN (UNII: 08232NY3SJ) HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK) MAGNESIUM STEARATE (UNII: 70097M6I30) MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) POLYDEXTROSE (UNII: VH2XOU12IE) POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E) SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2) | Ingredient Name | Strength |
| HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK) MAGNESIUM STEARATE (UNII: 70097M6I30) MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) POLYDEXTROSE (UNII: VH2XOU12IE) POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E) SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2) | SILICON DIOXIDE (UNII: ETJ7Z6XBU4) | |
| ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK) MAGNESIUM STEARATE (UNII: 70097M6I30) MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) POLYDEXTROSE (UNII: VH2XOU12IE) POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E) SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2) | STARCH, CORN (UNII: O8232NY3SJ) | |
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| POLYDEXTROSE (UNII: VH2XOU12IE) POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E) SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2) | MAGNESIUM STEARATE (UNII: 70097M6I30) | |
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| POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E) SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2) | POLYDEXTROSE (UNII: VH2XOU12IE) | |
| SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2) | POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) | |
| • • | POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E) | |
| STEADIC ACID (LINII) 4ELV/7765AD) | SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2) | |
| SILARIC ACID (UNII. 4LLV/200AF) | STEARIC ACID (UNII: 4ELV7Z65AP) | |
| TRIACETIN (UNII: XHX3C3X673) | TRIACETIN (UNII: XHX3C3X673) | |

| Product Characteristics | | | | |
|-------------------------|-------|--------------|----------|--|
| Color | white | Score | no score | |
| Shape | ROUND | Size | 10mm | |
| Flavor | | Imprint Code | 44;438 | |
| Contains | | | | |

| P | ackaging | | | |
|---|----------------------|--|-------------------------|-----------------------|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:0363-0438- 99 | 1 in 1 CARTON | 03/01/1999 | |
| 1 | | 100 in 1 BOTTLE; Type 0: Not a Combination Product | | |
| 2 | NDC:0363-0438- 12 | 1 in 1 CARTON | 03/01/1999 | 04/07/2019 |
| 2 | | 100 in 1 BOTTLE; Type 0: Not a Combination Product | | |
| 3 | NDC:0363-0438- 15 | 1 in 1 CARTON | 03/01/1999 | 10/08/2016 |
| 3 | | 50 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 | ANDA075139 | 03/01/1999 | | | |
| | | | | | |

Labeler - Walgreen Company (008965063)

| Establishment | | | |
|-------------------------|---------|-----------|----------------------------|
| Name | Address | ID/FEI | Business Operations |
| LNK International, Inc. | | 038154464 | pack(0363-0438) |

| Establishment | | | |
|-------------------------|---------|-----------|----------------------------|
| Name | Address | ID/FEI | Business Operations |
| LNK International, Inc. | | 832867837 | manufacture(0363-0438) |

| Establishment | | | |
|-------------------------|---------|-----------|----------------------------|
| Name | Address | ID/FEI | Business Operations |
| LNK International, Inc. | | 832867894 | manufacture(0363-0438) |

| Establishment | | | |
|-------------------------|---------|-----------|----------------------------|
| Name | Address | ID/FEI | Business Operations |
| LNK International, Inc. | | 967626305 | pack(0363-0438) |

| Establishment | | | |
|-------------------------|---------|-----------|----------------------------|
| Name | Address | ID/FEI | Business Operations |
| LNK International, Inc. | | 117025878 | manufacture(0363-0438) |

Revised: 5/2023 Walgreen Company