

IBUPROFEN- ibuprofen tablet, film coated RPK Pharmaceuticals, Inc.

IBUPROFEN 400 MG - 600 MG AND 800 MG TABLETS

ibuprofen tablets 400 mg - 600 mg- 800 mg medguide

Medication Guide for Nonsteroidal Anti-Inflammatory Drugs (NSAIDs)

What is the most important information I should know about medicines called Nonsteroidal Anti-Inflammatory Drugs (NSAIDs)?

NSAIDs can cause serious side effects, including:

- Increased risk of a heart attack or stroke that can lead to death. This risk may happen early in treatment and may increase:
 - with increasing doses of NSAIDs
 - with longer use of NSAIDs

Do not take NSAIDs right before or after a heart surgery called a “coronary artery bypass graft (CABG).”

Avoid taking NSAIDs after recent heart attack, unless your healthcare provider tells you to. You may have an increased risk of another heart attack if you take NSAIDs after a recent heart attack.

- Increased risk of bleeding, ulcers and tears (perforation) of the esophagus (tube leading from the mouth to the stomach), stomach and intestines:
 - anytime during use
 - without warning symptoms
 - that may cause death

The risk of getting an ulcer or bleeding increases with:

- past history of stomach ulcers, or stomach or intestinal bleeding with use of NSAIDs
- taking medicines called “corticosteroids”, “anticoagulants”, “SSRIs”, or “SNRIs”
- increasing doses of NSAIDs
- older age
- longer use of NSAIDs
- poor health
- smoking
- advanced liver disease
- drinking alcohol
- bleeding problems

NSAIDs should only be used:

- exactly as prescribed
- at the lowest dose possible for your treatment
- for the shortest time needed

What are NSAIDs?

NSAIDs are used to treat pain and redness, swelling, and heat (inflammation) from medical conditions such as different types of arthritis, menstrual cramps and other types of short-term pain.

Who should not take NSAIDs?

Do not take NSAIDs:

- if you have had an asthma attack, hives, or other allergic reaction with aspirin or any other NSAIDs
- right before or after heart bypass surgery.

Before taking NSAIDs tell your healthcare provider about all of your medical conditions, including if you:

- have liver or kidney problems
- if you have high blood pressure
- have asthma
- are pregnant or plan to become pregnant. Talk to your healthcare provider if you are considering taking NSAIDs during pregnancy. You should not take NSAIDs after 29 weeks of pregnancy.
- are breast feeding or plan to breast feed.

Tell your healthcare provider about all of the medicines you take, including prescription or over-the-counter medicines, vitamins or herbal supplements.

NSAIDs and some other medicines can interact with each other and cause serious side effects. Do not start taking any new medicine without talking to your healthcare provider first.

What are the possible side effects of NSAIDs?

NSAIDs can cause serious side effects, including:

See “What is the most important information I should know about medicines called Nonsteroidal Anti-inflammatory Drugs (NSAIDs)?”

- new or worse high blood pressure
- heart failure
- liver problems including liver failure
- kidney problems including kidney failure
- low red blood cells (anemia)
- life-threatening skin reactions
- life-threatening allergic reactions
- Other side effects of NSAIDs include: stomach pain, constipation, diarrhea, gas, heartburn, nausea, vomiting, and dizziness.

Get emergency help right away if you get any of the following symptoms:

- shortness of breath or trouble breathing
- chest pain
- weakness in one part or side of your body
- slurred speech
- swelling of the face or throat

Stop taking your NSAID and call your healthcare provider right away if you get any of the following symptoms:

- nausea
- more tired or weaker than usual
- diarrhea
- itching

- itching
- your skin or eyes look yellow
- indigestion or stomach pain
- flu-like symptoms
- vomit blood
- there is blood in your bowel movement or it is black and sticky like tar
- unusual weight gain
- skin rash or blisters with fever
- swelling of the arms, legs, hands and feet

If you take too much of your NSAID, Call your healthcare provider or get medical help right away.

These are not all the possible side effects of NSAIDs. For more information, ask your healthcare provider or pharmacist about NSAIDs.

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

Other information about NSAIDs

- Aspirin is an NSAID but it does not increase the chance of a heart attack. Aspirin can cause bleeding in the brain, stomach, and intestines. Aspirin can also cause ulcers in the stomach and intestines.
- Some NSAIDs are sold in lower doses without a prescription (over-the-counter). Talk to your healthcare provider before using over-the-counter NSAIDs for more than 10 days.

General information about the safe and effective use of NSAIDs

Medicines are sometimes prescribed for purposes other than those listed in a Medication Guide. Do not use NSAIDs for a condition for which it was not prescribed. Do not give NSAIDs to other people, even if they have the same symptoms that you have. It may harm them.

If you would like more information about NSAIDs, talk with your healthcare provider. You can ask your pharmacist or healthcare provider for information about NSAIDs that is written for health professionals.

This Medication Guide has been approved by the U.S. Food and Drug Administration.

Revision: R01/16

Manufactured for:
Time-Cap Labs, Inc.
7 Michael Avenue,
Farmingdale, NY 11735, USA

Manufactured by:
Marksans Pharma Ltd.
Plot No. L-82, L-83, Verna Indl. Estate,
Verna, Goa-403 722, India

HOW SUPPLIED

Product: 53002-3372

NDC: 53002-3372-1 15 TABLET, FILM COATED in a BOTTLE

NDC: 53002-3372-3 30 TABLET, FILM COATED in a BOTTLE

NDC: 53002-3372-5 50 TABLET, FILM COATED in a BOTTLE

Ibuprofen 400mg Tablets

NDC 53002-3372-3 30 TABLETS 30 TABLETS Rx# 213580111000 ORDER#131-38
 IBUPROFEN 400MG TABLETS, USP
 TMS-CAP LABS Generic for MOTRON 400MG
 LOT# 21334-042 EXP 02-28-2023
 LIST# 49483-602-01
 LOT# 21334-042
 EXP: 02-28-2023
 ORDER# 337-38
Rx only
 (Color Image - See patient instructions for details)

IBUPROFEN 400MG TABLETS
 TAKE 1 TABLET
 4 TIMES A DAY
 OR AS DIRECTED.

EACH FILM-COATED TABLET
 IBUPROFEN, USP 400MG
 DISPENSE A MEDICATION GUIDE
 PROVIDE TO EACH PATIENT

IMPORTANT: THIS DRUG IS A
 NONSTEROIDAL ANTI-INFLAMMATORY
 AGENT. TAKE WITH FOOD OR MILK.
 READ PATIENT MEDICATION CARD -
 FULLY BEFORE USING.

CLINIC NAME GOES HERE
 Patient Name _____
 Prescriber Name _____
 Date Dispensed: _____

LOT# 21334-042 EXP 02-28-2023
 Rx# 213580111000 FDA-3372
 30 ea IBUPROFEN 400MG TABLETS
 BILLING NDC# 49483-0602-01
 Rx# 213580111000
 30 ea IBUPROFEN 400MG TABLETS
 BILLING NDC# 49483-0602-01
 Rx# 213580111000
 30 ea IBUPROFEN 400MG TABLETS
 BILLING NDC# 49483-0602-01
 Rx# 213580111000
 30 ea IBUPROFEN 400MG TABLETS

IBUPROFEN 400MG TABLETS, USP 30 TABLETS
 DISCARD BY 02-28-2023
 NDC# 53002-3372-3 Rx#213580111000
 FEDERAL LAWS PROHIBIT REFILL OF THIS PRESCRIPTION WITHOUT THE PRESCRIBER'S HELP. IF YOU ARE A PATIENT OF THE REACH OF CHILDREN, CALL YOUR DOCTOR FOR MEDICAL ADVICE ABOUT THE EFFECTS OF THIS DRUG. REPORT SIDE EFFECTS TO FDA AT 1-800-FDA-1088
 Clinic Name Here
 PRESCRIBER NAME DATE
 PATIENT NAME

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:53002-3372(NDC:49483-602)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XY110QM) (IBUPROFEN - UNII:WK2XY110QM)	IBUPROFEN	400 mg

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
STARCH, CORN (UNII: O8232NY3SJ)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	13mm
Flavor		Imprint Code	121
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53002-3372-1	15 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2022	
2	NDC:53002-3372-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2022	
3	NDC:53002-3372-5	50 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090796	12/30/2015	

Labeler - RPK Pharmaceuticals, Inc. (147096275)

Establishment

Name	Address	ID/FEI	Business Operations
RPK Pharmaceuticals, Inc.		147096275	RELABEL(53002-3372) , REPACK(53002-3372)

Revised: 3/2022

RPK Pharmaceuticals, Inc.