



AGOG Pharma Ltd.



(WHO - GMP CERTIFIED - GOVT RECOGNISED EXPORT HOUSE)

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LEAFLET

IBUGO-400

(Ibuprofen Tablets BP 400 mg)

COMPOSITION :

Each film coated tablet contains:
Ibuprofen BP 400 mg
Excipients q.s.
Approved colour used.

INDICATIONS

Ibuprofen provides relief from mild to moderate pain in conditions such as dysmenorrhoea, migraine, post operative pain, rheumatic disorders such as ankylosing spondylitis, osteoarthritis and rheumatoid arthritis including juvenile rheumatoid arthritis and in other musculo-skeletal and joint disorders such as sprains and strains.

CONTRAINDICATIONS

Ibuprofen is contraindicated in the following conditions :

1. Bronchospasm
2. Known sensitivity to aspirin
3. Peptic ulcer
4. Recent gastrointestinal bleeding
5. Renal failure
6. Hypertension on treatment

PHARMACOLOGICAL ACTIONS

The propionic acid derivatives are all effective inhibitors of cyclo-oxygenase responsible for the biosynthesis of prostaglandins. All of these agents alter platelet function and prolong bleeding time.

Ibuprofen is well absorbed after oral administration. Ibuprofen is rapidly metabolised and eliminated in the urine. The excretion of Ibuprofen is virtually complete within 24 hours after the last dose. The serum half life is 1.8 to 2.0 hours.

WARNING

Bronchospasm may be precipitated in patients suffering from or with a previous history of bronchial asthma. Ibuprofen should not be given to patients in whom aspirin and other non-steroidal anti-inflammatory drugs induce the symptoms of asthma, rhinitis or urticaria.

PRECAUTIONS

Blurred or diminished vision have been reported. If a patient develops such complaints while receiving Ibuprofen tablets, the drug should be discontinued and the patient should have an ophthalmologic examination. Fluid retention and oedema have been reported in association with Ibuprofen; therefore, the drug should be used with caution in patients with a history of cardiac decompensation or hypertension. Ibuprofen should be used with caution in persons with intrinsic coagulation defects and those on anticoagulant therapy.

ADVERSE EFFECTS

The frequent adverse effects with Ibuprofen are gastrointestinal disturbances. Peptic ulceration and gastro-intestinal bleeding have been reported. Other side effects include dizziness, nervousness, skin rash, tinnitus, oedema, depression, blurred vision, drowsiness and other reactions are also common.

DOSAGE AND ADMINISTRATION

Initially 1200-1800 mg daily in 3-4 divided doses preferably after food; increase the dose if necessary to maximum 2400 mg daily; Maintenance dose of 600-1200 mg daily may be adequate. In case of juvenile rheumatoid arthritis, for children over 7 Kg bodyweight 30-40 mg/Kg daily in 3-4 divided doses. Fever and pain in children, Child over 7 Kg bodyweight 20-30 mg/Kg daily in divided doses, 3-7 years 100 mg 3-4 times daily, 8-12 years 200 mg 3-4 times daily.

STORAGE : Store under normal storage conditions (15°C to 30°C)

Protect from light.

Keep all medicines out of reach of children.

PRESENTATION: Blister pack of 10 x 10 Tablets.
Jar pack of 1000 Tablets.
Jar pack of 500 Tablets.



Manufactured in India by:

AGOG PHARMA LTD.

Plot No. 33, Sector II, The Vasai Taluka Indl. Co-op. Estate Ltd., Vasai (E), Dist. Thane. INDIA.

1195 / 01 / 01