125X155mm

KETOPROFEN

KETOPLAST

Rx **30 mg Plaster**
Non-Steroidal
Anti-Inflammatory Drug (NSAID)

FORMULATION:
Each plaster contains:
Ketoprofen…… 30 mg

DESCRIPTION:
Plaster of flesh-colored nonwoven fabric coated transparent and adhesive substance and covered its solid surface with a thin paper.

PHARMACODYNAMICS:
Topical application of Ketoprofen on the joints or muscles can increase the drug concentration at the target site as well as lower the concentration in the systemic circulation, thereby reducing stomach irritations and liver toxicity.

INDICATIONS:
Traumatic inflammation of muscles, sprains and strains. Relief of symptoms of muscular pain.

DOSAGE AND ADMINISTRATION:
Adults: Take off thin backing paper and apply to affected area twice daily, or upon advice of doctors.
Children: Not recommended as safety in children has not been established.

CONTRAINDICATIONS:
Ketoprofen plaster must not be used in patients with:
- hypersensitivity to this drug
- allergic reactions to any of the ingredient
- history of photosensitivity reactions
- third trimester of pregnancy
- pathological skin changes such as eczema; or in infected skin or open wounds

SPECIAL PRECAUTIONS:
- It is recommended to protect areas by wearing clothing during all the application of the plaster and two weeks following its discontinuation to avoid the risk of photosensitization.
- The plaster must not come into contact with mucous membrane or the eyes.
- The recommended length (1 week) should not be exceeded due to the risk of developing contact dermatitis and photosensitivity reactions which increases over time.
- Patients with asthma combined with chronic rhinitis, chronic sinusitis and/or nasal polyposis have a higher risk of allergy to aspirin and/or NSAIDs than the rest of the population.
- Should be a skin rash occur after plaster must be stopped. application, treatment must be stopped.
- Areas of skin treated with Ketoprofen plaster should not be exposed to direct sunlight or ultraviolet light, either during the treatment or for two weeks following treatment discontinuation, in order to avoid phototoxicity reactions and photoallergy.

PREGNANCY:
During the third trimester of pregnancy, all prostaglandin synthetase inhibitors including Ketoprofen may induce cardiopulmonary and renal toxicity in the fetus. At the end of the pregnancy, prolonged bleeding time in both mother and child may occur. Non-steroidal anti-inflammatory drugs may also delay labour. Therefore, Ketoprofen is contraindicated during the last trimester of pregnancy.

LACTATION:
As safety of this drug has not been established in lactation.

ADVERSE DRUG REACTION:
Occasionally redness, rashes, pruritus, dryness and irritation and rarely occasionally swelling occurs.
If these symptoms are serious, treatment with this preparation should be discontinued.

CAUTION:
Foods, Drugs, Devices and Cosmetics Act prohibits dispensing without prescription.

STORAGE CONDITION:
Store at temperatures not exceeding 30°C .

AVAILABILITY:
Paper and lint fabric sheet aluminum pack of 7plasters (Box of 20's)

For suspected adverse drug reaction, report to the FDA:www.fda.gov.ph. Patient should seek medical attention immediately at the first sign of any adverse drug reaction.

DRP-6567

Date of Renewal of Authorization: July 2021
Date of Revision of Package Insert: April 2023

Manufactured by:
Dae Hwa Pharmaceutical Co., Ltd.
495, Hanu-ro, Hoengseong-eup, Hoengseong-gun,
Gangwon-do, Republic of Korea

Imported by:
H & B Pharma International Inc.
21st Floor Unit 2102 The Podium West Tower 12 ADB Avenue,
Ortigas Center, Brgy. Wack-Wack Greenhills East,
Mandaluyong City, Metro Manila

Distributed by:
ONE PHARMA
One Pharma Marketing Inc.
L51 B21 Abel Nosce St., BF Resort Village,
Talon II, Las Piñas City, Metro Manila