

TRUECEPROL OD

For the use of a Registered Medical Practitioner or a Hospital or a Laboratory Only
Abbreviated Prescribing information for **Trueceprol OD [Oxaceprol 600mg sustained release tablets]**

[Please refer the complete prescribing information available at www.torrentpharma.com]

PHARMACOLOGICAL PROPERTIES: Oxaceprol prevents leukocyte infiltration into the joints, thus inhibiting an early step of inflammatory cascade and presenting a novel class of anti-inflammatory agents.

INDICATION: For the treatment of pain associated with osteoarthritis in adult patients.

DOSAGE AND ADMINISTRATION: The general recommended dose of the Oxaceprol SR Tablets 600 mg is one tablet once daily. Patients can safely use Oxaceprol SR Tablets 600 mg up to 42 days. Patient must consult the physician for further treatment with Oxaceprol SR Tablets 600 mg. Oxaceprol SR Tablets 600 mg should be taken before meals.

CONTRAINDICATION: It is contraindicated in patients with hypersensitivity to the active substance or to any of the excipients, pregnancy & Lactation, anticoagulant therapy with Vitamin K antagonists, hepatic impairment, and Renal impairment.

WARNINGS & PRECAUTIONS: Patients taking anticoagulants from the vitamin K antagonist group may not completely exclude the effect of Oxaceprol on blood coagulation. Therefore, when taking this medicine at the same time regular monitoring of prothrombin time is recommended. Patients can safely use Oxaceprol SR Tablets 600 mg up to 42 days. Patient must consult the physician for further treatment with Oxaceprol SR Tablets 600 mg.

DRUG INTERACTIONS: Patients taking anticoagulants from the vitamin K antagonist group may not completely exclude the effect of Oxaceprol on blood coagulation. Therefore, when taking Oxaceprol at the same time regular monitoring of prothrombin time is recommended.

ADVERSE REACTIONS: Nausea, diarrhea, dizziness and itching in skin.

MARKETED BY:

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PHARMA

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(Additional information is available on request)