FOUR STAGES OF OSTEOARTHRITIS (OA)

STAGE 0-1
PRE-OSTEOARTHRITIS
- At-risk dogs may appear asymptomatic or show subtle signs like “bunny hopping”
- Early diagnosis and education are important at this stage

STAGE 2
MILD
- Early clinical signs develop, but are easy to miss
- Begin therapy to manage stiffness, inflammation and pain
- Begin lifestyle changes to support healthier joints

STAGE 3
MODERATE
- At this stage, dogs have significant joint deterioration
- Therapeutic intervention is needed to manage pain and mobility

STAGE 4
SEVERE
- Untreated severe OA diminishes quality of life
- At this stage, therapeutic treatment may be the only way to end pain and improve mobility

EXERCISE RECOMMENDATION

NOTES:
PREVICOX® (firocoxib) is the easy-to-give, once-a-day way to help reduce pain and inflammation associated with osteoarthritis and certain types of surgery. Use the form below to track your dog's progress and share the results with your veterinarian.

Today's Date ___________________ Next Scheduled Appointment: ___________________

<table>
<thead>
<tr>
<th>WEEK</th>
<th>PREVICOX® DOSE</th>
<th>HOW IS YOUR DOG FEELING THIS WEEK? (lethargic, playful, etc.)</th>
<th>WHAT ACTIVITIES DID YOUR DOG DO THIS WEEK? (walk, extended play, stairs, etc.)</th>
<th>NOTES</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td></td>
<td>Lethargic</td>
<td>□ Walk  □ Extended Play</td>
<td>□ Stairs  □ Other</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Playful</td>
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<td>Other</td>
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<td>2</td>
<td></td>
<td>Lethargic</td>
<td>□ Walk  □ Extended Play</td>
<td>□ Stairs  □ Other</td>
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<td>Other</td>
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<td>3</td>
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<td>Lethargic</td>
<td>□ Walk  □ Extended Play</td>
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<td>4</td>
<td></td>
<td>Lethargic</td>
<td>□ Walk  □ Extended Play</td>
<td>□ Stairs  □ Other</td>
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</tr>
</tbody>
</table>

When you’re done, share this page with your veterinarian at your next appointment to ensure your pet is on the right course of treatment.

Important Safety Information
PREVICOX® (firocoxib) is for use in dogs only. People should not take PREVICOX. Keep PREVICOX and all medications out of the reach of children. PREVICOX, like other NSAIDs, may cause some side effects. Serious side effects associated with NSAID therapy in dogs can occur with or without warning, and, in rare situations, result in death. The most common side effects associated with PREVICOX therapy involve the digestive tract (vomiting and decreased food consumption). Liver and kidney problems have also been reported with NSAIDs. It is important to stop the medication and contact your veterinarian immediately if you think your dog has a medical problem or side effect while taking PREVICOX tablets. Evaluation for pre-existing conditions and regular monitoring are recommended for any pets on any medication, including PREVICOX. Use with other NSAIDs, corticosteroids or nephrotoxic medication should be avoided. Please refer to the prescribing information for complete details.

PREVICOX® is a registered trademark of Boehringer Ingelheim Animal Health USA Inc. ©2022 Boehringer Ingelheim Animal Health USA Inc., Duluth, GA. All rights reserved. US-PET-0255-2022.
Brief Summary: Before using PREVICOX, please consult the product insert, a summary of which follows:

Caution: Federal law restricts this drug to use by or on the order of a licensed veterinarian.

Indications: PREVICOX (firocoxib) Chewable Tablets are indicated for the control of pain and inflammation associated with osteoarthritis and for the control of postoperative pain and inflammation associated with soft tissue and orthopedic surgery in dogs.

Contraindications: Dogs with known hypersensitivity to firocoxib should not receive PREVICOX.

Warnings: Not for use in humans. Keep this and all medications out of the reach of children. Consult a physician in case of accidental ingestion by humans.

For oral use in dogs only. Use of this product at doses above the recommended 2.27 mg/lb (5.0 mg/kg) in pugs less than seven months of age has been associated with severe adverse reactions, including death (see Adverse Reactions). Due to tablet sizes and scoring, dogs weighing less than 12 lb (5.5 kg) cannot be accurately dosed. All dogs should undergo a thorough history and physical examination before the initiation of NSAID therapy. Appropriate laboratory testing to establish hematological and serum baseline data is recommended prior to and periodically during administration of any NSAID. Owners should be advised to observe for signs of potential drug toxicity (see Adverse Reactions and Animal Safety) and be given a Client Information Sheet about PREVICOX Chewable Tablets.

For technical assistance or to report suspected adverse events, call 1-888-637-4251. For animal information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS or www.fda.gov/reportanimalae.

Precautions: This product cannot be accurately dosed in dogs less than 12.5 pounds in body weight. Consider appropriate weight adjustments when switching from one NSAID to another or when switching from corticosteroid use to NSAID use.

As a class, cyclooxygenase inhibitor NSAIDs may be associated with renal, gastrointestinal and hepatic toxicity. Sensitivity to drug-associated adverse events varies with the individual patient. Dogs that have experienced adverse reactions from one NSAID may experience adverse reactions from another NSAID. Patients at greatest risk for select adverse reactions include those that are elderly, dehydrated, on concomitant diuretic therapy, or those with existing renal, cardiovascular, and/or hepatic dysfunction. Concomitant administration of potentially nephrotoxic drugs should be carefully approached and monitored. NSAIDs may inhibit the prostaglandins that maintain normal homeostatic function. Such anti-prostaglandin effects may result in clinically significant disease in patients with underlying or pre-existing disease that has not been previously diagnosed. Since NSAIDs possess the potential to produce gastrointestinal ulceration and/or gastrointestinal perforation, concomitant use of NSAIDs with other anti-inflammatory drugs such as NSAIDs, corticosteroids, and/or diuretics should be avoided. The concomitant use of NSAID and/or corticosteroids with other medicinal drugs may reduce the therapeutic effects of the latter drugs, or increase the toxicity of either drug. On the other hand, NSAIDs used concomitantly with anti-inflammatory drugs may reduce the toxicity of the latter drugs or increase the therapeutic effects of either drug. NSAIDs may reduce the efficacy of anticoagulants and antiplatelet agents by inhibiting the synthesis of thromboxane A2, a potent platelet aggregator. NSAIDs may reduce the therapeutic effectiveness of antihypertensive agents, particularly diuretics (see Drug Interactions).

Adverse Reactions Seen in Orthopedic Surgery:

Orthopedic Surgery:

Adverse Reactions Seen in U.S. Field Studies

Post-Approval Experience (Rev. 2020): The following adverse reactions are based on post-approval adverse drug event reports. The categories are listed in decreasing order of frequency by body system:

Gastrointestinal: Vomiting, anorexia, diarrhea, nausea, regurgitation, gastrointestinal perforation, haematemesis, haematochezia, weight loss, anorexia, vomiting, abdominal pain, ulceration, pruritus, diarrhea

Hepatic: Elevated ALT, elevated creatinine, polyuria, polydipsia, haematuria, urinary incontinence, proteinuria, kidney failure, ascites, urinary tract infection

Hematological: Neutropenia, Splenomegaly, Lymphadenopathy, petechiae, ecchymoses, lymphadenopathy, neutropenia, thrombocytopenia

Urinary: Elevated BUN, elevated creatinine, polyuria, polydipsia, haematuria, urinary incontinence, proteinuria, kidney failure, ascites

Respiratory: Tachypnea, dyspnea, tachycardia

Neurological/Behavioral: Seizures, pruritus, paresis, paresis, pain, salivation, tremors, ataxia, convulsions, diarrhea, agitation, ataxia, tremors, ataxia, hypothermia, hypothermia

Skin: Pruritus, alopecia, dermatitis, pruritus, dermatitis, pruritus, dermatitis, pruritus, dermatitis, pruritus

Other: Convulsions, salivation, vomiting, diarrhea, anorexia, weight loss, gastrointestinal ulceration, peritonitis, abdominal pain, hypersalivation, nausea

In some situations, death has been reported as an outcome of the adverse events listed above. For technical assistance or to report suspected adverse events, call 1-888-637-4251. For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS or www.fda.gov/reportanimalae.

Information for Patients: Owners of PREVICOX Chewable Tablets should be advised to monitor their dogs closely for signs of potential drug toxicity. Owners should be advised to notify their veterinarian if their dog develops any clinical signs consistent with drug toxicity. Owners should be advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets. Owners are advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets.

Information for Practitioners: Owners of PREVICOX Chewable Tablets should be advised of the potential for adverse reactions and be informed of the clinical signs associated with drug administration. Owners should be advised not to administer the drug to dogs with known hypersensitivity to firocoxib. Owners should be advised to observe the dog for signs of potential adverse reactions and be informed of the clinical signs associated with drug administration. Owners should be advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets. Owners are advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets.

Information for Vendors/Wholesalers/Drug Distributors: Owners of PREVICOX Chewable Tablets should be advised of the potential for adverse reactions and be informed of the clinical signs associated with drug administration. Owners should be advised not to administer the drug to dogs with known hypersensitivity to firocoxib. Owners should be advised to observe the dog for signs of potential adverse reactions and be informed of the clinical signs associated with drug administration. Owners should be advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets. Owners are advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets.

Information for Clinicians: Owners of PREVICOX Chewable Tablets should be advised of the potential for adverse reactions and be informed of the clinical signs associated with drug administration. Owners should be advised not to administer the drug to dogs with known hypersensitivity to firocoxib. Owners should be advised to observe the dog for signs of potential adverse reactions and be informed of the clinical signs associated with drug administration. Owners should be advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets. Owners are advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets.

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Information for Pet Owners: Owners of PREVICOX Chewable Tablets should be advised of the potential for adverse reactions and be informed of the clinical signs associated with drug administration. Owners should be advised not to administer the drug to dogs with known hypersensitivity to firocoxib. Owners should be advised to observe the dog for signs of potential adverse reactions and be informed of the clinical signs associated with drug administration. Owners should be advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets. Owners are advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets.

Information for Vets: Owners of PREVICOX Chewable Tablets should be advised of the potential for adverse reactions and be informed of the clinical signs associated with drug administration. Owners should be advised not to administer the drug to dogs with known hypersensitivity to firocoxib. Owners should be advised to observe the dog for signs of potential adverse reactions and be informed of the clinical signs associated with drug administration. Owners should be advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets. Owners are advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets.

Information for Laboratories: Owners of PREVICOX Chewable Tablets should be advised of the potential for adverse reactions and be informed of the clinical signs associated with drug administration. Owners should be advised not to administer the drug to dogs with known hypersensitivity to firocoxib. Owners should be advised to observe the dog for signs of potential adverse reactions and be informed of the clinical signs associated with drug administration. Owners should be advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets. Owners are advised to contact their veterinarian if they have any questions about the administration or side effects of PREVICOX Chewable Tablets.

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