

# Pyrocam 15 mg/ml oral suspension for pigs

Authorised

- Meloxicam

## Product identification

**Medicine name:**

Pyrocam 15 mg/ml oral suspension for pigs

---

**Active substance:**

Meloxicam

---

**Target species:**

Pig

---

**Route of administration:**

Oral use

---

## Product details

**Active substance and strength:**

Meloxicam

15.00 milligram(s) / 1.00 millilitre(s)

---

**Pharmaceutical form:**

Oral suspension

---

**Withdrawal period by route of administration:****Oral use:**

- 

**Pig**

- Meat and offal. 5 day

---

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QM01AC06

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Italy

---

**Package description:**

Cardboard box with HDPE bottle of 125 ml closed with a two-part tamper evident closure consisting of an outer white cap (PP), an internal natural colour screw closure (HDPE) and a mounted plug (LDPE) + plastic measuring syringe

Cardboard box with HDPE bottle of 250 ml closed with a two-part tamper evident closure consisting of an outer white cap (PP), an internal natural colour screw closure (HDPE) and a mounted plug (LDPE) + plastic measuring syringe

Cardboard box with HDPE bottle of 1000 ml closed with a two-part tamper evident closure consisting of an outer white cap (PP), an internal natural colour screw closure (HDPE) and a mounted plug (LDPE) + plastic measuring syringe

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Hybrid application – bioavailability studies cannot be used to demonstrate bioequivalence (Article 19(1)(b) of Regulation (EU) 2019/6)

---

**Marketing authorisation holder:**

HuVepharma

---

**Marketing authorisation date:**

9/07/2025

---

**Manufacturing sites for batch release:**

Biovet AD

---

**Responsible authority:**

Ministry Of Health

---

**Authorisation number:**

105708

---

**Date of authorisation status change:**

9/07/2025

---

**Reference member state:**

Belgium

---

**Procedure number:**

BE/V/0048/001

---

**Concerned member states:**

Austria Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France  
Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta  
Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain  
Sweden United Kingdom (Northern Ireland)

---

To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.