

# PYROCAM 20 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS AND HORSES

Authorised

- Meloxicam

## Product identification

**Medicine name:**

PYROCAM 20 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS AND HORSES

Pyrocum 20 mg/ml ενέσιμο διάλυμα για βοοειδή, χοίρους και άλογα

**Active substance:**

Meloxicam

**Target species:**

Pig

Cattle

Horse

**Route of administration:**

Intramuscular use

Subcutaneous use

Intravenous use

## Product details

### **Active substance and strength:**

Meloxicam

20.00 milligram(s) / 1.00 millilitre(s)

---

### **Pharmaceutical form:**

Solution for injection

---

### **Withdrawal period by route of administration:**

#### **Intramuscular use:**

- 

##### **Pig**

- Meat and offal. 5 day

#### **Subcutaneous use:**

- 

##### **Cattle**

- Meat and offal. 15 day
- Milk. 5 day

#### **Intravenous use:**

- 

##### **Cattle**

- Meat and offal. 15 day
- Milk. 5 day

- 

##### **Horse**

- Meat and offal. 5 day
- Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

---

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

QM01AC06

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Greece

---

**Package description:**

Cardboard box with 1 colourless glass injection vial type I containing 20 ml and closed with a bromobutyl rubber stopper and sealed with an aluminium cap.

Cardboard box with 1 colourless glass injection vial type I containing 50 ml and closed with a bromobutyl rubber stopper and sealed with an aluminium cap.

Cardboard box with 1 colourless glass injection vial type I containing 100 ml and closed with a bromobutyl rubber stopper and sealed with an aluminium cap.

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Generic application (Article 18 of Regulation (EU) 2019/6)

---

**Marketing authorisation holder:**

HuVepharma

---

**Marketing authorisation date:**

16/12/2024

---

**Manufacturing sites for batch release:**

Biovet AD

---

**Responsible authority:**

National Organization For Medicines

---

**Authorisation number:**

151515/17-12-2024/K-0255102

**Date of authorisation status change:**

16/12/2024

**Reference member state:**

France

**Procedure number:**

FR/V/0471/001

**Concerned member states:**

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

**Generic of:**

600000001532

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

## Documents

Summary of Product Characteristics

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

Package Leaflet

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