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

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:
Slovenia

Available in:
Slovenia

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:
20/08/2024

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0793/002

Date of authorisation status change:
20/08/2024

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

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.

Labelling

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

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