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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:

Denmark

Available in:

Denmark

Package description:

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 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 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:

3/06/2024

Manufacturing sites for batch release:

Biovet AD

Responsible authority:

Danish Medicines Agency

Authorisation number:

69166

Date of authorisation status change:

3/06/2024

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)

To consult adverse reactions on veterinary medicinal products please go to
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