

Melosolute 5 mg/ml Solution for injection for cattle, pigs, dogs and cats

Authorised

- Meloxicam

Product identification

Medicine name:

Melosolute 5 mg/ml Solution for injection for cattle, pigs, dogs and cats

Active substance:

Meloxicam

Target species:

Cattle

Pig

Dog

Cat

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Meloxicam

5.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. 15 day

-

Pig

- Meat and offal. 5 day

Intravenous use:

-

Cattle

- Meat and offal. 15 day

-

Pig

- Meat and offal. 5 day

Subcutaneous use:

-

Cattle

- Meat and offal. 15 day

-

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:

Germany

Available in:

Germany

Package description:

10 ml injection vial colourless glass (type I), closed with a bromobutyl rubber stopper and sealed with an aluminium cap in a cardboard box

100 ml injection vial colourless glass (type I), closed with a bromobutyl rubber stopper and sealed with an aluminium cap in a cardboard box

20 ml injection vial colourless glass (type I), closed with a bromobutyl rubber stopper and sealed with an aluminium cap in a cardboard box

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:

11/05/2012

Manufacturing sites for batch release:

CP-Pharma Handelsgesellschaft mbH

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

401591.00.00

Date of authorisation status change:

9/11/2017

Reference member state:

Netherlands

Procedure number:

NL/V/0164/001

Concerned member states:

France Germany Hungary

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.

Combined File of all Documents

Final PuAR Melosolute NL_V_0164_001-002_DC.docx.pdf