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Inflacam 20 mg/ml - Solution for injection

Authorised

Meloxicam

Product identification

Medicine name:

Inflacam 20 mg/ml - Solution for injection

Active substance:

Meloxicam

Target species:

Cattle

Horse

Pig

Route of administration:

Intramuscular use Intravenous use Subcutaneous use

Product details

Active substance and strength:

Meloxicam 20.00 milligram(s) / 1.00 Vial

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

•

Cattle

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

•

Horse

- Meat and offal. 5 day 5 days

•

Pig

- Meat and offal. 5 day 5 days

Intravenous use:

•

Cattle

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

•

Horse

- Meat and offal. 5 day 5 days

•

Pig

- Meat and offal. 5 day 5 days

Subcutaneous use:

Cattle

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

•

Horse

- Meat and offal. 5 day 5 days

•

Pig

- Meat and offal. 5 day 5 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

OM01AC06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

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

Package description:

Packaging:Vial (glass), Package_size:1 vial, Content:250 ml

Packaging:Vial (glass), Package_size:1 vial, Content:100 ml

Packaging:Vial (glass), Package_size:1 vial, Content:50 ml

Packaging:Vial (glass), Package_size:1 vial, Content:20 ml

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:

Chanelle Pharmaceuticals Manufacturing Ltd

Marketing authorisation date:

9/12/2011

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited Eurovet Animal Health BV Labiana Life Sciences S.A.

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

9/12/2011

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents
English (PDF) Published on: 5/12/2024 <u>Download</u>
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