Metaxx 5 mg/ml solution for injection for cattle, pigs, dogs and cats

Authorised

- Meloxicam

Product identification

Medicine name:
Metaxx 5 mg/ml solution for injection for cattle, pigs, dogs and cats

Active substance:
Meloxicam

Target species:
Dog
Cat
Cattle
Pig

Route of administration:
Subcutaneous use
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Meloxicam
5.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Concentrate for solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Dog**
  - Not applicable. no withdrawal period

- **Cat**
  - Not applicable. no withdrawal period

- **Cattle**
  - Meat and offal. no withdrawal period 15 days

Intramuscular use:

- **Pig**
  - Meat and offal. no withdrawal period 5 days

Intravenous use:

- **Cattle**
  - Meat and offal. no withdrawal period

15 days Not authorised for use in animals 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

Package description:
Colourless glass (type I) injection vial , closed with a rubber stopper and sealed with an aluminium cap
Colourless glass (type I) injection vial , closed with a rubber stopper and sealed with an aluminium cap. Cardboard box of 1 vial of 50 ml
Colourless glass (type I) injection vial , closed with a rubber stopper and sealed with an aluminium cap. Cardboard box of 1 vial of 250 ml
Colourless glass (type I) injection vial , closed with a rubber stopper and sealed with an aluminium cap. Cardboard box of 1 vial of 100 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic (abridged application) - art 13(1)

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
20/12/2022

Manufacturing sites for batch release:
Produlab Pharma B.V.
Alfasan Nederland B.V.

Responsible authority:
Danish Medicines Agency

Authorisation number:
66776

Date of authorisation status change:
20/12/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0375/001

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

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Documents

Combined File of all Documents

English (PDF)
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Summary of Product Characteristics

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