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:
Latvia

Package description:
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian

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:
30/11/2022

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

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/DCP/22/0046

Date of authorisation status change:
30/11/2022

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)

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: 17/05/2023
Updated on: 13/03/2026
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