

Animeloxan, 20 mg/ml, solution for injection for cattle, pigs and horses

Authorised

- Meloxicam

Product identification

Medicine name:

Animeloxan, 20 mg/ml, solution for injection for cattle, pigs and horses

Active substance:

Meloxicam

Target species:

Cattle

Horse

Pig

Route of administration:

Intravenous use

Subcutaneous use

Intramuscular 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:

Intravenous use:

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Cattle

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

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Horse

- Milk. no withdrawal period

Do not use in horses producing milk for human consumption.

- Meat and offal. 5 day

Subcutaneous use:

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Cattle

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

Intramuscular use:

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Pig

- Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AC06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Available in:

Greece

Package description:

(ID4) 1200 millilitre(s): unspecified outer container with 12 Vial (Glass) each with 100 millilitre(s)

(ID3) 600 millilitre(s): unspecified outer container with 12 Vial (Glass) each with 50 millilitre(s)

(ID2) 100 millilitre(s): unspecified outer container with 1 Vial with 100 millilitre(s)

(ID1) 50 millilitre(s): unspecified outer container with 1 Vial with 50 millilitre(s)

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:

aniMedica GmbH

Marketing authorisation date:

5/12/2019

Manufacturing sites for batch release:

aniMedica GmbH

Industrial Veterinaria S.A.

aniMedica Herstellungs GmbH

Responsible authority:

National Organization For Medicines

Authorisation number:

17585/14-02-2025/K-0239001

Date of authorisation status change:

13/02/2025

Reference member state:

Germany

Procedure number:

DE/V/0311/002

Concerned member states:

Austria Bulgaria Cyprus Czechia Estonia Greece Hungary Ireland Italy
Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Spain
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 14/02/2022

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Combined File of all Documents

Package Leaflet

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