

ALGENAMIC 40 mg/ml solution for injection for cattle, pigs, dogs and cats

Authorised

- Tolfenamic acid

Product identification

Medicine name:

ALGENAMIC 40 mg/ml solution for injection for cattle, pigs, dogs and cats

Active substance:

Tolfenamic acid

Target species:

Cattle

Pig

Dog

Cat

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Tolfenamic acid

40.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. no withdrawal period

Meat and offal: 12 days intramuscular use / 4 days intravenous use

- Milk. no withdrawal period

Milk: Zero days intramuscular use/ 24 hours intravenous use

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Pig

- Meat and offal. 16 day

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Cattle

- Meat and offal. no withdrawal period

Meat and offal: 12 days intramuscular use / 4 days intravenous use

- Milk. no withdrawal period

Milk: Zero days intramuscular use/ 24 hours intravenous use

Intravenous use:

-

Cattle

- Meat and offal. no withdrawal period

Meat and offal: 12 days intramuscular use / 4 days intravenous use

- Milk. no withdrawal period

Milk: Zero days intramuscular use/ 24 hours intravenous use

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Cattle

- Meat and offal. no withdrawal period

Meat and offal: 12 days intramuscular use / 4 days intravenous use

- Milk. no withdrawal period

Milk: Zero days intramuscular use/ 24 hours intravenous use

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AG02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

Cardboard box with 15 glass vials of 250 ml

Cardboard box with 10 glass vials of 100 ml

Cardboard box with 5 glass vial of 20 ml

Cardboard box with 1 glass vial of 250 ml

Cardboard box with 1 glass vial of 100 ml

Cardboard box with 1 glass vial of 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:

Vetpharma Animal Health S.L.

Marketing authorisation date:

27/08/2021

Manufacturing sites for batch release:

Mevet S.A.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/3959714 4/2021

Date of authorisation status change:

27/08/2021

Reference member state:

Spain

Procedure number:

ES/V/0382/001

Concerned member states:

Austria Belgium Croatia Cyprus France Germany Greece Italy Poland
Portugal Romania

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 22/12/2023

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Package Leaflet

Labelling

Package Leaflet and Labelling

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