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KETOPROFEN Bioveta 100 mg/ml solution for injection

Authorised

Ketoprofen

Product identification

Medicine name:

KETOPROFEN Bioveta 100 mg/ml solution for injection KETOPROFEN Bioveta 100 mg/ml Injektionslösung für Rinder, Pferde und Schweine

Active substance:

Ketoprofen

Target species:

Cattle

Horse

Pig

Route of administration:

Intravenous use

Intramuscular use

Product details

Active substance and strength:

Ketoprofen

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

•

Cattle

- Meat and offal. 1 day
- Milk. 0 hour

•

Horse

- Meat and offal. 1 day
- Milk. no withdrawal period

Nepoužívat u koní, jejichž mléko je určeno pro lidskou spotřebu.,

Intramuscular use:

•

Cattle

- Meat and offal. 4 day
- Milk. 0 hour

•

Pig

- Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AE03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Package description:

glass vial 1 x 100.0. 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:

Bioveta a.s.

Marketing authorisation date:

12/04/2024

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

842117

Date of authorisation status change:

12/04/2024

Reference member state:

Czechia

Procedure number:

CZ/V/0183/001

Concerned member states:

Austria Bulgaria France Greece Hungary Poland Romania Slovakia

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

Documents

Summary of Product Characteristics

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Labelling

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