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KoniFlun 83 mg/ml Injektionslösung für Rinder, Pferde und Schweine

Authorised

- Flunixin meglumine

Product identification

Medicine name:
KoniFlun 83 mg/ml Injektionslösung für Rinder, Pferde und Schweine

Active substance:
Flunixin meglumine

Target species:
Cattle
Horse
Pig

Route of administration:
Intravenous use
Intramuscular use

Product details

Active substance and strength:
Flunixin meglumine

83.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Cattle**
  - Milk. 24 hour
  - Meat and offal. 4 day
- **Horse**
  - Milk. no withdrawal period

Nicht bei Tieren anwenden, deren Milch für den menschlichen Verzehr vorgesehen ist.

  - Meat and offal. 5 day

Intramuscular use:

- **Pig**
  - Meat and offal. 24 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AG90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
Available only in German
Available only in German
Available only in German

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:
Konivet GmbH

Marketing authorisation date:
12/08/2019

Manufacturing sites for batch release:
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402606.00.00

Date of authorisation status change:
12/08/2019

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

Documents

Combined File of all Documents

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