Belamox 200 mg/ml Pulver und Lösungsmittel zur Herstellung einer Injektionslösung für Rinder, Pferde, Schweine

Authorised

- Amoxicillin sodium

Product identification

Medicine name:
Belamox 200 mg/ml Pulver und Lösungsmittel zur Herstellung einer Injektionslösung für Rinder, Pferde, Schweine

Active substance:
Amoxicillin sodium

Target species:
Cattle
Cattle (calf)
Horse
Pig

Route of administration:
Intravenous use
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Amoxicillin sodium
213.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Powder and solvent for solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Cattle**
  - Milk. 1 day
  - Meat and offal. 5 day
- **Cattle (calf)**
  - Meat and offal. 5 day
- **Horse**
  - Milk. 1 day
  - Meat and offal. 5 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 9 day
  - Milk. 72 hour
- **Cattle (calf)**
  - Meat and offal. 9 day
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Horse

- Meat and offal. 16 day
- Milk. 72 hour

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Pig

- Meat and offal. 9 day

Intramuscular use:

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Cattle

- Meat and offal. 9 day
- Milk. 72 hour

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Cattle (calf)

- Meat and offal. 9 day

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Horse

- Meat and offal. 16 day
- Milk. 72 hour

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Pig

- Meat and offal. 9 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

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Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
Available only in German
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Bela-Pharm GmbH & Co. KG

Marketing authorisation date:
19/02/1999

Manufacturing sites for batch release:
Bela-Pharm GmbH & Co. KG

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
6932991.00.00

Date of authorisation status change:
28/08/2006

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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