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Multimox pulver og solvens til injektionsvæske, opløsning 212 mg/ml

Authorised

- Amoxicillin sodium

Product identification

Medicine name:
Multimox pulver og solvens til injektionsvæske, opløsning 212 mg/ml

Active substance:
Amoxicillin sodium

Target species:
Pig
Cattle

Route of administration:
Intramuscular use
Intravenous use

Product details

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

Pharmaceutical form:
Powder and solvent for solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Meat and offal. 9 day
- **Cattle**
  - Meat and offal. 9 day
  - Milk. 3 day

Intravenous use:

- **Cattle**
  - Meat and offal. 5 day
  - Milk. 24 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:
Denmark

Available in:
Denmark

Package description:
Available only in Danish
Available only in Danish

Additional information

Entitlement type:
Marketing Authorisation

Marketing authorisation holder:
Scanvet Animal Health A/S

Marketing authorisation date:
2/11/2017

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

Responsible authority:
Danish Medicines Agency

Authorisation number:
58257

Date of authorisation status change:
2/11/2017

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

Documents

Summary of Product Characteristics

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