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Amoxicilline 100 inj.,suspensie voor injectie

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Amoxicilline 100 inj.,suspensie voor injectie

Active substance:
Amoxicillin trihydrate

Target species:
Cattle
Pig
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Amoxicillin trihydrate
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Milk. 84 hour
  - Meat and offal. 28 day
- **Pig**
  - Meat and offal. 28 day

Subcutaneous use:

- **Cattle**
  - Milk. 84 hour
  - Meat and offal. 28 day
- **Pig**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Available only in Dutch

Available only in Dutch
Available only in Dutch
Available only in Dutch

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Dopharma Research B.V.

Marketing authorisation date:
16/01/1992

Manufacturing sites for batch release:
Dopharma International B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 2275

Date of authorisation status change:
8/02/2016

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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