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Synulox RTU 140/35 mg/ml Injektionssuspension für Rinder, Schweine (Ferkel, Mastschweine), Hunde und Katzen

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:
Synulox RTU 140/35 mg/ml Injektionssuspension für Rinder, Schweine (Ferkel, Mastschweine), Hunde und Katzen

Active substance:
Amoxicillin trihydrate
Potassium clavulanate

Target species:
Cattle
Dog
Cat
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

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

Potassium clavulanate
41.65 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. 4 day
  - Meat and offal. 14 day
- **Pig**
  - Meat and offal. 16 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

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:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Deutschland GmbH

Marketing authorisation date:
17/11/2003

Manufacturing sites for batch release:
Haupt Pharma Latina S.r.l.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
400606.00.00

Date of authorisation status change:
9/07/2009

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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