SYNULOX RTU 140/35 mg/ml injekčná suspenzia

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:
SYNULOX RTU 140/35 mg/ml injekčná suspenzia

Active substance:
Amoxicillin trihydrate
Potassium clavulanate

Target species:
Cattle
Pig
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Amoxicillin trihydrate
160.73 milligram(s) / 1.00 millilitre(s)

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

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 42 day
  - Milk. 60 hour
- **Pig**
  - Meat and offal. 31 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 42 day
  - Milk. 60 hour

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:
Slovakia

Available in:

Slovakia

Package description:
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Zoetis Ceska Republika s.r.o.

Marketing authorisation date:
3/08/1999

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

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/082/99-S

Date of authorisation status change:
3/08/1999

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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