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Synulox RTU suspensija injekcijām liellopiem, cūkām, suņiem un kaķiem

Authorised

- Amoxicillin
- Potassium clavulanate

Product identification

Medicine name:
Synulox RTU suspensija injekcijām liellopiem, cūkām, suņiem un kaķiem

Active substance:
Amoxicillin

Potassium clavulanate

Target species:
Dog
Cat
Cattle
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

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

Potassium clavulanate
35.00 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

60 stundas (5 slaukšanas reizes, ja govs tiek slaukta divas reizes dienā)

- **Pig**
  - Meat and offal. 31 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:
Latvia

Available in:

Latvia

Package description:
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
28/06/2002

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

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/02/1476

Date of authorisation status change:
30/06/2002

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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