SYNULOX RTU ενέσιμο εναιώρημα για βοοειδή, χοίρους, σκύλους και γάτες

Authorised

- Potassium clavulanate
- Amoxicillin

Product identification

Medicine name:
SYNULOX RTU ενέσιμο εναιώρημα για βοοειδή, χοίρους, σκύλους και γάτες

Active substance:
Potassium clavulanate

Amoxicillin

Target species:
Cattle
Pig
Cat
Dog

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Potassium clavulanate
35.00 milligram(s) / 1.00 millilitre(s)

Amoxicillin
140.00 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
- **Pig**
  - Meat and offal. 26 day
- **Cat**
  - Not applicable. no withdrawal period
- **Dog**
  - Not applicable. no withdrawal period

Subcutaneous use:

- **Cat**
  - Not applicable. no withdrawal period
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Dog

- Not applicable. no withdrawal period

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

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

Authorisation status:
Valid

Authorised in:
Greece

Available in:
Greece

Package description:
Available only in Greek
Available only in Greek

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
Zoetis Hellas S.A.

Marketing authorisation date:
7/06/1999

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

Responsible authority:
National Organization For Medicines

Authorisation number:
31804/09-04-2021/K-0077403

Date of authorisation status change:
8/04/2021

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