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Synulox Ready to Use Injection 140 mg/ml + 35 mg/ml suspension for injection

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
Synulox Ready to Use Injection 140 mg/ml + 35 mg/ml suspension for injection

Active substance:
Potassium clavulanate

Amoxicillin trihydrate

Target species:
Cattle
Dog
Cat
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

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

Amoxicillin trihydrate
160.70 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. 80 hour
- **Pig**
  - Meat and offal. 31 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 42 day
  - Milk. 80 hour
- **Pig**
  - Meat and offal. 31 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

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

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
Type III glass vials of 100 ml containing an off-white sterile, non-aqueous suspension. The vials are sealed with a rubber bungand an aluminium seal. Each vial contains a sterile off-white to pale buff coloured, smooth fluid, readily dispersable suspension.
Type III glass vials of 40 ml containing an off-white sterile, non-aqueous suspension. The vials are sealed with a rubber bungand an aluminium seal. Each vial contains a sterile off-white to pale buff coloured, smooth fluid, readily dispersable suspension.

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 Belgium S.A.

Marketing authorisation date:
2/05/2014

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10387/075/001

Date of authorisation status change:
2/05/2014

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

Documents

Summary of Product Characteristics