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SYNULOX 140 mg/ml + 35 mg/ml suspensión inyectable

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:
SYNULOX 140 mg/ml + 35 mg/ml suspensión inyectable

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
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. 54 day

- **Pig**
  - Meat and offal. 27 day

- **Cattle**
  - Milk. 108 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

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

Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
Available only in Spanish

Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Spain S.L.

Marketing authorisation date:
15/07/1992

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
450 ESP

Date of authorisation status change:
15/07/1992

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

Documents

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.