SYNULOX suspensão injetável para suínos, bovinos, cães e gatos

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
SYNULOX suspensão injetável para suínos, bovinos, cães e gatos

Active substance:
Potassium clavulanate
Amoxicillin trihydrate

Target species:
Cattle
Pig
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

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

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 42 day
  - Milk. 2 day
- **Pig**
  - Meat and offal. 42 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

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

Authorisation status:
Valid

Authorised in:
Portugal

Package description:
Available only in Portuguese
Available only in Portuguese
Available only in Portuguese
Available only in Portuguese
Available only in Portuguese

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 Portugal Lda.

Marketing authorisation date:
21/05/1997

Manufacturing sites for batch release:
ZOETIS MANUFACTURING ITALIA S.r.l.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
51178

Date of authorisation status change:
1/09/2006

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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