AMPIDEXALONE SUSPENSION INJECTABLE POUR BOVINS EQUINS ET PORCINS

Authorised

- Ampicillin trihydrate
- Dexamethasone
- COLISTIN SULFATE

Product identification

Medicine name:
AMPIDEXALONE SUSPENSION INJECTABLE POUR BOVINS EQUINS ET PORCINS

Active substance:
Ampicillin trihydrate
Dexamethasone
COLISTIN SULFATE

Target species:
Cattle
Pig
Equid

Route of administration:
Intramuscular use
Subcutaneous use
Intraperitoneal use

Product details

Active substance and strength:

Ampicillin trihydrate
100.46 milligram(s) / 1.00 millilitre(s)

Dexamethasone
0.25 milligram(s) / 1.00 millilitre(s)

COLISTIN SULFATE
0.25 million international units / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 21 day
  - Milk. 5 day
- **Pig**
  - Meat and offal. 21 day
- **Equid**
  - Meat and offal. 21 day
  - Milk. 5 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 21 day
  - Milk. 5 day

- **Pig**
  - Meat and offal. 21 day
- **Equid**
  - Meat and offal. 21 day
  - Milk. 5 day

Intraperitoneal use:

- **Cattle**
  - Meat and offal. 21 day
  - Milk. 5 day
- **Pig**
  - Meat and offal. 21 day
- **Equid**
  - Meat and offal. 21 day
  - Milk. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01RV01

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

Authorisation status:
Valid

Authorised in:
France

Package description:
Available only in French

Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Dopharma France S.A.S.

Marketing authorisation date:
26/10/1989

Manufacturing sites for batch release:
Dopharma France

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/1053151 8/1989

Date of authorisation status change:
26/10/2009

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 and Labelling

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