FERCOBSANG

Authorised

This information is not available for this product.

Product identification

Medicine name:
FERCOBSANG

Active substance:
This information is not available for this product.

Target species:
Cattle
Pig
Horse
Horse (mare)
Sheep
Goat
Cat
Dog

Route of administration:
Intramuscular use
Subcutaneous use
Oral use

Product details

Active substance and strength:
This information is not available for this product.

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Horse (mare)**
  - Milk. 0 day
- **Sheep**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day

Subcutaneous use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Horse (mare)**
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day

- Milk. 0 day

- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day

Oral use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Cat**
- **Horse**
  - Meat and offal. 0 day
- **Horse (mare)**
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QB03AE04

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

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Authorisation status:
Valid

Authorised in:
France

Package description:
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:
Vetoquinol S.A.

Marketing authorisation date:
30/06/1992

Manufacturing sites for batch release:
Vetoquinol

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

Authorisation number:
FR/V/5599387 0/1992

Date of authorisation status change:
30/06/2012

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

Documents

Summary of Product Characteristics

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