CORULON, liofilizzato e solvente per soluzione iniettabile per bovini, equini, ovini, caprini, suini, cani e gatti

Authorised

- Chorionic gonadotrophin

Product identification

Medicine name:
CORULON, liofilizzato e solvente per soluzione iniettabile per bovini, equini, ovini, caprini, suini, cani e gatti

Active substance:
Chorionic gonadotrophin

Target species:
Cattle
Horse
Sheep
Goat
Dog
Cat
Pig

Route of administration:
Intravenous use
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Chorionic gonadotrophin
1500.00 international unit(s) / 1.00 Bottle

Pharmaceutical form:
Lyophilisate and solvent for solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Dog**
- **Cat**

- **Pig**
  - Meat and offal. 0 day

Intramuscular use:

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

Subcutaneous use:

- **Cattle**

- Meat and offal. 0 day
- Milk. 0 hour

- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour

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

- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour

- **Pig**
  - Meat and offal. 0 day

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG03GA01

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

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Available only in Italian
Available only in Italian

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:
Intervet International B.V.

Marketing authorisation date:
4/02/1982

Manufacturing sites for batch release:
INTERVET INTERNATIONAL B.V.
Intervet International GmbH

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
1/01/2009

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

Documents

Combined File of all Documents

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