

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

-

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

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

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

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