Facilpart 10 ui/ml solução injetável

Authorised

- Oxytocin

Product identification

Medicine name:
Facilpart 10 ui/ml solução injetável

Active substance:
Oxytocin

Target species:
Cattle
Goat
Horse
Sheep
Pig
Dog
Cat

Route of administration:
Subcutaneous use
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Oxytocin
10.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

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

Intramuscular use:

-

Cattle

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

•

Goat

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

•

Horse

- Meat and offal. 0 day

•

Sheep

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

•

Pig

- Meat and offal. 0 day

Intravenous use:

•

Cattle

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

•

Goat

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

•

Horse

- Meat and offal. 0 day

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH01BB02

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

Authorisation status:
Valid

Authorised in:
Portugal

Available 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:
Iapsa Portuguesa Pecuaria Lda.

Marketing authorisation date:
8/07/1994

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
51076

Date of authorisation status change:
1/12/2016

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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