PARTOVET 10 UI/ml SOLUCION INYECTABLE

Authorised

- Oxytocin

Product identification

Medicine name:
PARTOVET 10 UI/ml SOLUCION INYECTABLE

Active substance:
Oxytocin

Target species:
Horse (mare)
Dog (bitch)
Cat (adult female)
Goat (adult female)
Cattle (cow)
Pig (female)
Sheep (ewe)

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous 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:
Intramuscular use:

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

Intravenous use:

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

Subcutaneous use:

- **Horse (mare)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat (adult female)**
  - Meat and offal. 0 day
  - Milk. 0 day
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Cattle (cow)

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

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Pig (female)

- Meat and offal. 0 day

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Sheep (ewe)

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH01BB02

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

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Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Divasa Farmavic S.A.

Marketing authorisation date:
15/07/1992

Manufacturing sites for batch release:
Divasa Farmavic S.A.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
449 ESP

Date of authorisation status change:
15/07/1992

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

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Labelling

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