PARTOVET 10 IU/ml injekčný roztok

Authorised

- OXYTOCIN SYNTHETIC

Product identification

Medicine name:
PARTOVET 10 IU/ml injekčný roztok

Active substance:
OXYTOCIN SYNTHETIC

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

Route of administration:
Intramuscular use
Subcutaneous use
Intravenous use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle (cow)**
  - All relevant tissues. 0 day — All relevant tissues: Zero days
- **Horse (mare)**
  - All relevant tissues. 0 day — All relevant tissues: Zero days
- **Pig (sow)**
  - All relevant tissues. 0 day — All relevant tissues: Zero days
- **Goat (adult female)**
  - All relevant tissues. 0 day — All relevant tissues: Zero days
- **Sheep (ewe)**
  - All relevant tissues. 0 day — All relevant tissues: Zero days

Subcutaneous use:

- **Cattle (cow)**

- All relevant tissues. 0 day All relevant tissues: Zero days

- **Horse (mare)**
  - All relevant tissues. 0 day All relevant tissues: Zero days

- **Pig (sow)**
  - All relevant tissues. 0 day All relevant tissues: Zero days

- **Goat (adult female)**
  - All relevant tissues. 0 day All relevant tissues: Zero days

- **Sheep (ewe)**
  - All relevant tissues. 0 day All relevant tissues: Zero days

Intravenous use:

- **Cattle (cow)**
  - All relevant tissues. 0 day Zero days

- **Horse (mare)**
  - All relevant tissues. 0 day Zero days

- **Pig (sow)**
  - All relevant tissues. 0 day Zero days

- **Goat (adult female)**
  - All relevant tissues. 0 day Zero days

- **Sheep (ewe)**
  - All relevant tissues. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH01BB02

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

Authorisation status:
Valid

Authorised in:
Slovakia

Available in:
Slovakia

Package description:
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Divasa Farmavic S.A.

Marketing authorisation date:
30/04/2001

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

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/080/04-S

Date of authorisation status change:
30/04/2001

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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