Partovet (10 ui/ml de oxitocina sintética), solução injetável para bovinos, equinos, suínos, caprinos, ovinos, cães e gatos

Authorised

- OXYTOCIN SYNTHETIC

Product identification

Medicine name:
Partovet (10 ui/ml de oxitocina sintética), solução injetável para bovinos, equinos, suínos, caprinos, ovinos, cães e gatos

Active substance:
OXYTOCIN SYNTHETIC

Target species:
Cattle
Goat
Horse
Sheep
Pig
Dog
Cat

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

Subcutaneous use:

- **Cattle**

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

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

- **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

Package description:
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:
Divasa Farmavic S.A.

Marketing authorisation date:
11/02/1988

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

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:

287/01/10NFVPT

Date of authorisation status change:
1/04/2026

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.