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OXYTOCIN BIO, 5IU/ml, Injekční roztok

Authorised

- Oxytocin

Product identification

Medicine name:
OXYTOCIN BIO, 5IU/ml, Injekční roztok

Active substance:
Oxytocin

Target species:
Cattle (cow)
Horse (mare)
Sheep
Goat
Pig (sow)
Dog (bitch)

Route of administration:
Intravenous use
Subcutaneous use
Intramuscular use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Horse (mare)**
  - 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 (sow)**
  - Meat and offal. 0 day

Subcutaneous use:

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Horse (mare)**
  - 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 (sow)**
  - Meat and offal. 0 day

Intramuscular use:

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Horse (mare)**
  - Meat and offal. 0 day
  - Milk. 0 hour
-

Sheep

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

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Goat

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

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Pig (sow)

- Meat and offal. 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:
Czechia

Package description:
Available only in Czech
Available only in Czech
Available only in Czech

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:
Bioveta a.s.

Marketing authorisation date:
12/06/2008

Manufacturing sites for batch release:
Bioveta a.s.

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

Authorisation number:
96/032/08-C

Date of authorisation status change:
22/05/2013

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

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.