

OXYTOCIN BIOVETA 5 UI/ ml solutie injectabila pentru vaci, iepe, oi, capre, scroafe si catele

Authorised

- Oxytocin

Product identification

Medicine name:

OXYTOCIN BIOVETA 5 UI/ ml solutie injectabila pentru vaci, iepe, oi, capre, scroafe si catele

Active substance:

Oxytocin

Target species:

Cattle (cow)
Horse (mare)
Sheep
Goat (adult female)
Pig (sow)
Dog (bitch)

Route of administration:

Intravenous use
Intramuscular use
Subcutaneous 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 day

-

Horse (mare)

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

-

Sheep

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

-

Goat (adult female)

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

-

Pig (sow)

- Meat and offal. 0 day

Intramuscular use:

-

Cattle (cow)

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

-

Horse (mare)

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

-

Sheep

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

-

Goat (adult female)

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

-

Pig (sow)

- Meat and offal. 0 day

Subcutaneous use:

-

Cattle (cow)

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

-

Sheep

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

-

Horse (mare)

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

-

Goat (adult female)

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

-

Pig (sow)

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01BB02

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Romania

Package description:

Available only in Romanian

Available only in Romanian

Available only in Romanian

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 Romania S.R.L.

Marketing authorisation date:

6/02/2014

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

190044

Date of authorisation status change:

19/06/2025

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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