

Oxytocina Pituitaria Calier 20 IU/ml šķīdums injekcijām liellopiem, zirgiem, aitām, kazām, cūkām, suņiem un kaķiem

Not
authorised

- Oxytocin

Product identification

Medicine name:

Oxytocina Pituitaria Calier 20 IU/ml šķīdums injekcijām liellopiem, zirgiem, aitām, kazām, cūkām, suņiem un kaķiem

Active substance:

Oxytocin

Target species:

Pig
Dog
Cat
Goat
Horse
Cattle
Sheep

Route of administration:

Intravenous use

Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Oxytocin

20.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

-

Pig

- Meat and offal. 2 day

-

Goat

- Milk. 2 day

- Meat and offal. 2 day

-

Horse

- Milk. 2 day

- Meat and offal. 2 day

-

Cattle

- Milk. 2 day

- Meat and offal. 2 day

-

Sheep

- Milk. 2 day

- Meat and offal. 2 day

Intramuscular use:

-

Pig

- Meat and offal. 2 day

-

Sheep

- Milk. 2 day
- Meat and offal. 2 day

-

Cattle

- Milk. 2 day
- Meat and offal. 2 day

-

Horse

- Meat and offal. 2 day
- Milk. 2 day

-

Goat

- Meat and offal. 2 day
- Milk. 2 day

Subcutaneous use:

-

Pig

- Meat and offal. 2 day

-

Cattle

- Meat and offal. 2 day
- Milk. 2 day

-

Horse

- Meat and offal. 2 day
- Milk. 2 day

-

Sheep

- Meat and offal. 2 day
- Milk. 2 day

-

Goat

- Meat and offal. 2 day
- Milk. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01BB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Latvia

Package description:

Available only in Latvian

Available only in Latvian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:

Laboratorios Calier S.A.

Marketing authorisation date:

16/02/2001

Manufacturing sites for batch release:

Laboratorios Calier S.A.

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/NRP/01/1291

Date of authorisation status change:

5/06/2025

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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

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.