

ZOPITUINA

Not authorised

- Oxytocin

Product identification

Medicine name:

ZOPITUINA

Active substance:

Oxytocin

Target species:

Horse (mare)

Dog (bitch)

Cat (adult female)

Goat (adult female)

Cattle (cow)

Pig (female)

Sheep (ewe)

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Oxytocin

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

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

•

Horse (mare)

- Meat and offal. 0 day

- Milk. 0 day

•

Goat (adult female)

- Meat and offal. 0 day

- Milk. 0 day

•

Cattle (cow)

- Meat and offal. 0 day

- Milk. 0 day

•

Pig (female)

- Meat and offal. 0 day

•

Sheep (ewe)

- Meat and offal. 0 day

- Milk. 0 day

Intravenous use:

•

Horse (mare)

- Meat and offal. 0 day

- Milk. 0 day

•

Goat (adult female)

- Meat and offal. 0 day

- Milk. 0 day

•

Cattle (cow)

- Meat and offal. 0 day

- Milk. 0 day

•

Pig (female)

- Meat and offal. 0 day

•

Sheep (ewe)

- Meat and offal. 0 day

- Milk. 0 day

Subcutaneous use:

•

Horse (mare)

- Meat and offal. 0 day

- Milk. 0 day

•

Goat (adult female)

- Meat and offal. 0 day

- Milk. 0 day

•

Cattle (cow)

- Meat and offal. 0 day

- Milk. 0 day

-

Pig (female)

- Meat and offal. 0 day

-

Sheep (ewe)

- Meat and offal. 0 day

- Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01BB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Spain

Package description:

Available only in [Spanish](#)

Available only in [Spanish](#)

Available only in [Spanish](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Laboratorios E Industrias Iven S.A.

Marketing authorisation date:

15/07/1992

Manufacturing sites for batch release:

Laboratorios E Industrias Iven S.A.

Responsible authority:

Spanish Agency For Medicines And Health Products

Authorisation number:

444 ESP

Date of authorisation status change:

31/10/2023

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.