

LABIPITUIN, 10 IU/ML ενέσιμο διάλυμα για αγελάδες, φοράδες, προβατίνες, αίγες, σύες, θηλυκούς σκύλους και γάτες

Authorised

- OXYTOCIN SYNTHETIC

Product identification

Medicine name:

LABIPITUIN, 10 IU/ML ενέσιμο διάλυμα για αγελάδες, φοράδες, προβατίνες, αίγες, σύες, θηλυκούς σκύλους και γάτες

Active substance:

OXYTOCIN SYNTHETIC

Target species:

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

Route of administration:

Intramuscular and intravenous use

Subcutaneous 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 and intravenous use:

-

Horse (mare)

- Meat and offal. 3 day
- Milk. 24 hour

-

Cattle (cow)

- Meat and offal. 3 day
- Milk. 24 hour

-

Sheep (ewe)

- Meat and offal. 3 day
- Milk. 24 day

-

Goat (adult female)

- Meat and offal. 3 day
- Milk. 24 hour

-

Pig (female)

- Meat and offal. 3 day

-

Dog (bitch)

- Meat and offal. no withdrawal period Δεν εφαρμόζεται

-

Cat

- Meat and offal. no withdrawal period Δεν εφαρμόζεται

Subcutaneous use:

-

Horse (mare)

- Meat and offal. 3 day
- Milk. 24 hour

-

Cattle (cow)

- Meat and offal. 3 day
- Milk. 24 hour

-

Sheep (ewe)

- Meat and offal. 3 day
- Milk. 24 hour

-

Goat (adult female)

- Meat and offal. 3 day
- Milk. 24 hour

-

Pig (female)

- Meat and offal. 3 day

-

Dog (bitch)

- Meat and offal. no withdrawal period Δεν εφαρμόζεται

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Cat

- Meat and offal. no withdrawal period Δεν εφαρμόζεται

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01BB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Package description:

Available only in Greek

Available only in Greek

Available only in Greek

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone)

Marketing authorisation holder:

Chellafarm Vet A.E.

Marketing authorisation date:

28/02/2005

Manufacturing sites for batch release:

Labiana Life Sciences S.A.

Responsible authority:

National Organization For Medicines

Authorisation number:

95628/22-10-2021/K-0114701

Date of authorisation status change:

21/10/2021

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

Documents

Summary of Product Characteristics

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