

LIDOR 20 MG/ML SOLUTION FOR INJECTION FOR HORSES, DOGS AND CATS

Authorised

- Lidocaine hydrochloride monohydrate

Product identification

Medicine name:

LIDOR 20 MG/ML SOLUTION FOR INJECTION FOR HORSES, DOGS AND CATS

Active substance:

Lidocaine hydrochloride monohydrate

Target species:

Dog

Cat

Horse

Route of administration:

Epidural use

Subcutaneous use

Perineural use

Ocular use

Intraarticular use

Product details

Active substance and strength:

Lidocaine hydrochloride monohydrate

24.65 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Epidural use:

-

Horse

- Meat and offal. 3 day

- Milk. 3 day

Subcutaneous use:

-

Horse

- Meat and offal. 3 day

- Milk. 3 day

Perineural use:

-

Horse

- Meat and offal. 3 day

- Milk. 3 day

Ocular use:

-

Horse

- Meat and offal. 3 day

- Milk. 3 day

Intraarticular use:

-

Horse

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN01BB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Available in:

Spain

Package description:

Cardbord box with 50 ml
Cardbord box with 250 ml
Cardbord box with 5 x 100 ml
Cardbord box with 100 ml
Cardbord box with 5 x 50 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Vetviva Richter GmbH

Marketing authorisation date:

16/02/2018

Manufacturing sites for batch release:

Vetviva Richter GmbH

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

3623 ESP

Date of authorisation status change:

16/02/2018

Reference member state:

France

Procedure number:

FR/V/0318/001

Concerned member states:

Austria Belgium Czechia Denmark Estonia Finland Germany Iceland Italy
Latvia Lithuania Netherlands Poland Portugal Spain Sweden

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.

Combined File of all Documents

eu-puar-frv0318001-mr-rpe340-en.pdf