

# 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

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**Active substance:**

Lidocaine hydrochloride monohydrate

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**Target species:**

Dog  
Cat  
Horse

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**Route of administration:**

Epidural use  
Subcutaneous use  
Perineural use  
Ocular use  
Intraarticular use

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## Product details

### **Active substance and strength:**

Lidocaine hydrochloride monohydrate

24.65 milligram(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Solution for injection

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### **Withdrawal period by route of administration:**

#### **Epidural use:**

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##### **Horse**

- Meat and offal. 3 day

- Milk. 3 day

#### **Subcutaneous use:**

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##### **Horse**

- Meat and offal. 3 day

- Milk. 3 day

#### **Perineural use:**

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##### **Horse**

- Meat and offal. 3 day

- Milk. 3 day

#### **Ocular use:**

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##### **Horse**

- Meat and offal. 3 day

- Milk. 3 day

**Intraarticular use:**

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**Horse**

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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN01BB02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Czechia

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Vetviva Richter GmbH

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**Marketing authorisation date:**

16/03/2018

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**Manufacturing sites for batch release:**

Vetviva Richter GmbH

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/011/18-C

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**Date of authorisation status change:**

16/03/2018

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**Reference member state:**

France

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**Procedure number:**

FR/V/0318/001

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**Concerned member states:**

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

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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