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



• Lidocaine hydrochloride monohydrate

## Product identification

#### **Medicine name:**

LIDOR 20 MG/ML SOLUTION FOR INJECTION FOR HORSES, DOGS AND CATS Lidor vet. 20 mg/ml Injektionsvätska, lösning

#### **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:**

•

Dog

•

Cat

•

#### Horse

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

## **Subcutaneous use:**

•

Dog

•

Cat

•

#### Horse

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

#### **Perineural use:**

•

Dog

.

Cat

Horse	
- Meat and offal. 3 day	
- Milk. 3 day	
Ocular use:	
•	
Dog	
•	
Cat	
•	
Horse	
- Meat and offal. 3 day	
- Milk. 3 day	
Intraarticular use:	
•	
Dog	
•	
Cat	
•	
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:	
Sweden	

#### **Available in:**

Sweden

## 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:

12/01/2018

## Manufacturing sites for batch release:

Vetviva Richter GmbH

### Responsible authority:

**Swedish Medical Products Agency** 

#### **Authorisation number:**

56081

### Date of authorisation status change:

12/01/2018

#### 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**

Package Leaflet

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

Summary of Product Characteristics

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