File downloaded on 2026-11-28
Source URL: https://medicines.health.europa.eu/veterinary/en/600000031114

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:
Germany

Available in:
Germany

Package description:
Cardbord box with 5 x 50 ml
Cardbord box with 100 ml
Cardbord box with 5 x 100 ml
Cardbord box with 250 ml
Cardbord box with 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:
25/01/2018

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402425.00.00

Date of authorisation status change:
25/01/2018

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

Combined File of all Documents

English (PDF)

Published on: 10/10/2025
Updated on: 13/03/2026
Download

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