

Mepidor 20 mg/ml solution for injection for horses

Authorised

- Mepivacaine hydrochloride

Product identification

Medicine name:

Mepidor 20 mg/ml solution for injection for horses

Active substance:

Mepivacaine hydrochloride

Target species:

Horse

Route of administration:

Epidural use
Intraarticular use

Product details

Active substance and strength:

Mepivacaine hydrochloride
20.00 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. 72 hour

Intraarticular use:

-

Horse

- Meat and offal. 3 day
- Milk. 72 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN01BB03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

Cardboard box with clear glass vial type I, bromobutyl rubber stopper or bromobutyl stopper with a fluorinated polymer coating and aluminium capPack sizes: 10 ml.

Cardboard box with clear glass vial type I, bromobutyl rubber stopper or bromobutyl stopper with a fluorinated polymer coating and aluminium capPack sizes: 5 x 10 ml.

Cardboard box with clear glass vial type I, bromobutyl rubber stopper or bromobutyl stopper with a fluorinated polymer coating and aluminium capPack sizes: 6 x 10 ml.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Vetviva Richter GmbH

Marketing authorisation date:

9/11/2017

Manufacturing sites for batch release:

Vetviva Richter GmbH

Responsible authority:

Danish Medicines Agency

Authorisation number:

59052

Date of authorisation status change:

9/11/2017

Reference member state:

Ireland

Procedure number:

IE/V/0425/001

Concerned member states:

Austria Denmark Estonia Finland France Germany Netherlands Spain
Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 23/11/2025

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Combined File of all Documents