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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

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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: 6 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: 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: 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
Updated on: 14/03/2026
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Combined File of all Documents