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Mepiblock 20 mg/ml solution for injection for horses

Authorised

- Mepivacaine hydrochloride

Product identification

Medicine name:
Mepiblock 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. 2 day
  - Milk. 2 day

Intraarticular use:

- **Horse**
  - Meat and offal. 2 day
  - Milk. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01BB03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Norway

Available in:
Norway

Package description:
Clear type I glass vial containing 10 ml, with a red chlorobutyl rubber stopper and aluminium flip cap, available in carton of six.

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:
Dechra Regulatory B.V.

Marketing authorisation date:
4/12/2017

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
16-11375

Date of authorisation status change:
4/12/2017

Reference member state:
Ireland

Procedure number:
IE/V/0375/001

Concerned member states:
Austria Belgium Denmark Finland France Germany Italy Netherlands Norway Poland Portugal Spain Sweden

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 14/06/2026
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Package Leaflet

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Combined File of all Documents