

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

Mepiblock Vet. 20 mg/ml injektionsvæske, opløsning

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:

Denmark

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:

9/11/2017

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Danish Medicines Agency

Authorisation number:

58633

Date of authorisation status change:

9/11/2017

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)

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