Mepidor 20 mg/ml Solution for Injection

Authorised

- Mepivacaine hydrochloride
- Mepivacaine hydrochloride

Product identification

Medicine name:
Mepidor 20 mg/ml solution for injection
Mepidor 20 mg/ml Solution for Injection

Active substance:
Mepivacaine hydrochloride
Mepivacaine hydrochloride

Target species:
Horse (non food-producing)
Horse

Route of administration:
Infiltration
Epidural use
Intraarticular use

Product details

Active substance and strength:
Mepivacaine hydrochloride

20.00 milligram(s) / 1.00 millilitre(s)

Mepivacaine hydrochloride
20.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Infiltration:

- **Horse (non food-producing)**

Epidural use:

- **Horse**

Intraarticular use:

- **Horse**

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:
United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

Package description:
Cardboard box with clear glass vials type I, 6X10 ml, bromobutyl rubber stopper or bromobutyl stopper with a fluorinated polymer coating and aluminium cap.
Cardboard box with clear glass vials type I, 10 ml, bromobutyl rubber stopper or bromobutyl stopper with a fluorinated polymer coating and aluminium cap.

Cardboard box with clear glass vials type I, 5X10 ml, bromobutyl rubber stopper or bromobutyl stopper with a fluorinated polymer coating and aluminium cap.

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:
4/10/2016

Manufacturing sites for batch release:
Richter Pharma AG

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 22080/4008

Date of authorisation status change:
4/10/2016

Reference member state:
Portugal

Procedure number:
PT/V/0128/001

Concerned member states:
Belgium Iceland Italy Norway United Kingdom (Northern Ireland)

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

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