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Sedivet 10 mg/ml Oplossing voor injectie

Authorised

- Romifidine hydrochloride

Product identification

Medicine name:
Sedivet 10 mg/ml Oplossing voor injectie

Active substance:
Romifidine hydrochloride

Target species:
Horse

Route of administration:
Intravenous use

Product details

Active substance and strength:
Romifidine hydrochloride
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Horse**
  - Meat and offal. 6 day
  - Milk. no withdrawal period

Do not use in animals producing milk for human consumption

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN05CM93

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

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Sedivet 20 ml Vial Solution for injection

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
31/12/1991

Manufacturing sites for batch release:
Labiana Life Sciences S.A.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V156676

Date of authorisation status change:
16/06/2022

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

Documents

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.