Sedivet 10 mg/ml oplossing voor injectie voor paarden

Authorised

- Romifidine hydrochloride

Product identification

Medicine name:
Sedivet 10 mg/ml oplossing voor injectie voor paarden

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN05CM93

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Available only in Dutch

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
21/05/1992

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

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 7232

Date of authorisation status change:
26/04/2018

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

Documents

Combined File of all Documents

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