Sedivet vet. 10 mg/ml injeksjonsvæske, oppløsning til hest

Authorised

- Romifidine hydrochloride

Product identification

Medicine name:
Sedivet vet. 10 mg/ml injeksjonsvæske, oppløsning til hest

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. 10 day

Not permitted for use in mares 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:
Norway

Available in:
Norway

Package description:
Available only in Norwegian

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:
19/10/1993

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

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
7913

Date of authorisation status change:
19/10/1993

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

Documents

Package Leaflet

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

Summary of Product Characteristics

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