Sedivet vet. 10 mg/ml Injektionsvätska, lösning

Authorised

- Romifidine hydrochloride

Product identification

Medicine name:
Sedivet vet. 10 mg/ml Injektionsvätska, lösning

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. 10 day

Ej tillåtet för användning till lakterande djur som producerar mjölk för human konsumtion.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN05CM93

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

Authorisation status:
Valid

Authorised in:
Sweden

Available in:
Sweden

Package description:
Available only in Swedish

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:
5/02/1993

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

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
11753

Date of authorisation status change:
5/02/1993

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.