

Sedivet Vet. injektionsvæske, opløsning 10 mg/ml

Authorised

- Romifidine hydrochloride

Product identification

Medicine name:

Sedivet Vet. injektionsvæske, opløsning 10 mg/ml

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05CM93

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

Available only in Danish

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:

26/07/1995

Manufacturing sites for batch release:

Labiana Life Sciences S.A.

Responsible authority:

Danish Medicines Agency

Authorisation number:

14896

Date of authorisation status change:

26/07/1995

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

Documents

Summary of Product Characteristics

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