

# 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

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**Active substance:**

Romifidine hydrochloride

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**Target species:**

Horse

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**Route of administration:**

Intravenous use

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## Product details

**Active substance and strength:**

Romifidine hydrochloride

10.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Intravenous use:**

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**Horse**

- Meat. 10 day

Not permitted for use in mares producing milk for human consumption

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN05CM93

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Norway

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**Available in:**

Norway

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**Package description:**

Available only in Norwegian

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Boehringer Ingelheim Vetmedica GmbH

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**Marketing authorisation date:**

19/10/1993

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**Manufacturing sites for batch release:**

Labiana Life Sciences S.A.

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**Responsible authority:**

Norwegian Medical Products Agency

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**Authorisation number:**

7913

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**Date of authorisation status change:**

19/10/1993

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://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.