

# SEDIVET SOLUTION INJECTABLE

Authorised

- Romifidine hydrochloride

## Product identification

**Medicine name:**

SEDIVET SOLUTION INJECTABLE

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**Active substance:**

Romifidine hydrochloride

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**Target species:**

Horse (mare)

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 (mare)**

- Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine.

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**Horse**

- Meat and offal. 6 day

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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:**

France

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**Available in:**

France

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**Package description:**

Available only in French

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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 Animal Health France

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**Marketing authorisation date:**

28/06/1995

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**Manufacturing sites for batch release:**

Labiana Life Sciences S.A.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/1717685 1/1995

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**Date of authorisation status change:**

28/06/2010

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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

Summary of Product Characteristics

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

Package Leaflet and Labelling

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