

Sedivet 10 mg/ml Solution for Injection for horses

Authorised

- Romifidine hydrochloride

Product identification

Medicine name:

Sedivet 10 mg/ml Solution for Injection for horses

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:**

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Horse

- Meat and offal. 6 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05CM93

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

20 ml clear Type I PhEur glass vial with grey bromobutyl stopper containing a clearcolourless aqueous solution.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

29/03/1996

Manufacturing sites for batch release:

Labiana Life Sciences S.A.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10454/013/001

Date of authorisation status change:

29/03/1996

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

Documents

Summary of Product Characteristics