SEDIVET 10 mg/ml SOLUCION INYECTABLE PARA CABALLOS

Authorised

- Romifidine hydrochloride

Product identification

Medicine name:
SEDIVET 10 mg/ml SOLUCION INYECTABLE PARA CABALLOS

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. 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:
Spain

Available in:
Spain

Package description:
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
28/06/1996

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
1107 ESP

Date of authorisation status change:
28/06/1996

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

Documents

Labelling

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Summary of Product Characteristics

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Package Leaflet

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