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Sedivet

Authorised

- Romifidine hydrochloride

Product identification

Medicine name:
Sedivet

Active substance:
Romifidine hydrochloride

Target species:
Horse

Route of administration:
Intravenous use

Product details

Active substance and strength:
Romifidine hydrochloride
0.20 gram(s) / 20.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Horse**
  - Meat and offal. 6 day
  - Milk. no withdrawal period

Nicht bei Stuten anwenden, deren Milch für den menschlichen Verzehr vorgesehen ist.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN05CM93

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
Available only in German

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:
13/09/1995

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

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
23715.00.00

Date of authorisation status change:
28/10/2005

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

Documents

Combined File of all Documents

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