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Sedan, 10mg/ml, Injekční roztok

Authorised

- Acepromazine

Product identification

Medicine name:
Sedan, 10mg/ml, Injekční roztok

Active substance:
Acepromazine

Target species:
Horse

Route of administration:
Intravenous use
Intramuscular use

Product details

Active substance and strength:
Acepromazine
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. no withdrawal period

O ošetření musí být proveden zápis do průkazu koně.,

  - Milk. no withdrawal period

O ošetření musí být proveden zápis do průkazu koně.,

Intramuscular use:

- **Horse**
  - Meat and offal. no withdrawal period

O ošetření musí být proveden zápis do průkazu koně.,

  - Milk. no withdrawal period

O ošetření musí být proveden zápis do průkazu koně.,

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN05AA04

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

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Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech
Available only in Czech

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bioveta a.s.

Marketing authorisation date:
23/10/2019

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/080/19-C

Date of authorisation status change:
28/01/2022

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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