

SEDALIN gel, 35mg/ml, Perorální gel

Authorised

- Acepromazine

Product identification

Medicine name:

SEDALIN gel, 35mg/ml, Perorální gel

Active substance:

Acepromazine

Target species:

Dog

Horse

Route of administration:

Oral use

Product details

Active substance and strength:

Acepromazine

35.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral gel

Withdrawal period by route of administration:**Oral use:**

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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

Authorisation status:

Valid

Authorised in:

Czechia

Available in:

Czechia

Package description:

Available only in Czech

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

Vetoquinol s.r.o.

Marketing authorisation date:

29/12/1997

Manufacturing sites for batch release:

Vetoquinol Biowet Sp. z o.o.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/1341/97-C

Date of authorisation status change:

26/08/2011

To consult adverse reactions on veterinary medicinal products please go to 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

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.