

Sedan, 35mg/ml, Oral gel

Authorised

- Acepromazine

Product identification

Medicine name:

Sedan, 35mg/ml, Oral gel

Active substance:

Acepromazine

Target species:

Horse

Dog

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:

-

Horse

- Meat and offal. no withdrawal period

The treatment must be recorded into the horse's passport.,

- Milk. no withdrawal period

The treatment must be recorded into the horse's passport.,

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05AA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

Plastic Applicator 1 x 10.0 millilitre(s)

Plastic Applicator 1 x 1.0 millilitre(s)

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:

24/06/2022

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/026/MR/22-S

Date of authorisation status change:

24/06/2022

Reference member state:

Czechia

Procedure number:

CZ/V/0174/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Denmark Estonia France Germany
Greece Hungary Italy Latvia Lithuania Netherlands Norway Poland Portugal
Romania Slovakia Spain Sweden

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

Documents

Combined File of all Documents

English (PDF)

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