Solupam 5 mg/ml solution for injection for dogs and cats

Authorised

- Diazepam

Product identification

Medicine name:
Solupam 5 mg/ml solution for injection for dogs and cats
Solupam 5 mg/ml raztopina za injiciranje za pse in mačke

Active substance:
Diazepam

Target species:
Dog
Cat

Route of administration:
Intravenous use

Product details

Active substance and strength:
Diazepam
5.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05BA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovenia

Package description:

Box with 1 vial of 50 ml
Box with 1 vial of 5 ml
Box with 1 vial of 20 ml
Box with 1 vial of 10 ml
Multi-pack with 6 boxes each containing 1 vial of 5 mL
Multi-pack with 6 boxes each containing 1 vial of 20 mL
Multi-pack with 6 boxes each containing 1 vial of 10 mL
Multi-pack with 10 boxes each containing 1 vial of 5 mL
Multi-pack with 10 boxes each containing 1 vial of 20 mL
Multi-pack with 10 boxes each containing 1 vial of 10 mL

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:
Dechra Regulatory B.V.

Marketing authorisation date:
15/01/2019

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0648/001

Date of authorisation status change:
15/01/2019

Reference member state:
Netherlands

Procedure number:
NL/V/0242/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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.

Combined File of all Documents

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.
