Diazedor 5 mg/ml Injektionslösung für Hunde und Katzen

Authorised

- Diazepam
- Diazepam

Product identification

Medicine name:
Diazedor 5 mg/ml Injektionslösung für Hunde und Katzen

Active substance:
Diazepam
Diazepam

Target species:
Dog
Cat
Dog
Cat

Route of administration:
Intravenous use
Intravenous use

Product details

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

Diazepam
5.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN05BA01

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

Authorisation status:
Valid

Authorised in:
Sweden

Available in:
Sweden

Package description:
Colourless glass ampoules, type I, with 2 ml solution for injection, closed with a chlorobutyl rubber stopper and either an aluminium pull off cap or an aluminium/plastic flip off cap. Pack size: 10 x 2 ml in a cardboard box.
Colourless glass ampoules, type I, with 2 ml solution for injection, closed with a chlorobutyl rubber stopper and either an aluminium pull off cap or an aluminium/plastic flip off cap. Pack size: 5 x 2 ml in a cardboard box.
Colourless glass vials, type I, with 10 ml solution for injection, closed with a chlorobutyl rubber stopper and either an aluminium pull off cap or an aluminium/plastic flip off cap. Pack size: 1 x 10 ml in a cardboard box.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Vetviva Richter GmbH

Marketing authorisation date:
15/03/2018

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
56075

Date of authorisation status change:
15/03/2018

Reference member state:
Austria

Procedure number:
AT/V/0017/001

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

Generic of:

600000004401

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

Documents

Package Leaflet

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Summary of Product Characteristics

English (PDF)
Published on: 13/03/2025
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Labelling

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Combined File of all Documents

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