

Domosedan 10 mg/ml solution for injection for horses and cattle

Authorised

- DETOMIDINE HYDROCHLORIDE FOR VETERINARY USE

Product identification

Medicine name:

Domosedan 10 mg/ml solution for injection for horses and cattle

Active substance:

DETOMIDINE HYDROCHLORIDE FOR VETERINARY USE

Target species:

Cattle

Horse

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

DETOMIDINE HYDROCHLORIDE FOR VETERINARY USE

10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. 2 day
- Milk. 12 hour

-

Horse

- Meat and offal. 2 day

Intravenous use:

-

Cattle

- Meat and offal. 2 day
- Milk. 12 hour

-

Horse

- Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05CM90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Package description:

1 x Type I clear glass vial containing 20 ml, closed with a chlorobutyl rubber stopper and an aluminium overseal, in a cardboard box.

1 x Type I clear glass vial containing 5 ml, closed with a chlorobutyl rubber stopper and an aluminium overseal, in a cardboard box.

6 x Type I clear glass vials containing 5 ml, closed with a chlorobutyl rubber stopper and an aluminium overseal, in a cardboard box.

10 x Type I clear glass vials containing 5 ml, closed with a chlorobutyl rubber stopper and an aluminium overseal, in a cardboard box.

5 x Type I clear glass vials containing 20 ml, closed with a chlorobutyl rubber stopper and an aluminium overseal, in a cardboard box.

10 x Type I clear glass vials containing 20 ml, closed with a chlorobutyl rubber stopper and an aluminium overseal, in a cardboard box.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Orion Corporation

Marketing authorisation date:

4/08/1998

Manufacturing sites for batch release:

Orion Corporation

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 2973

Date of authorisation status change:

5/06/2024

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

Documents

Combined File of all Documents

English (PDF)

Published on: 12/12/2025

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