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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
  - Milk. 12 hour

Intravenous use:

- **Cattle**
  - Meat and offal. 2 day
  - Milk. 12 hour
- **Horse**
  - Meat and offal. 2 day
  - Milk. 12 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05CM90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Norway

Package description:
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.
1 x Type I clear glass vial containing 5 ml, closed with a chlorobutyl rubber stopper and an aluminium overseal, in a cardboard box.
1 x Type I clear glass vial 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:
6/05/1985

Manufacturing sites for batch release:
Orion Corporation

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
7006

Date of authorisation status change:

6/05/2010

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.

Combined File of all Documents