

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

Authorised

- DETOMIDINE HYDROCHLORIDE FOR VETERINARY USE

## Product identification

### **Medicine name:**

Domosedan vet. 10 mg/ml Injektionsvätska, lösning

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

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Cattle**

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

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**Horse**

- Meat and offal. 2 day

**Intravenous use:**

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**Cattle**

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

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**Horse**

- Meat and offal. 2 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN05CM90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Sweden

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Orion Corporation

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**Marketing authorisation date:**

20/06/1985

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**Manufacturing sites for batch release:**

Orion Corporation

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**Responsible authority:**

Swedish Medical Products Agency

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**Authorisation number:**

10243

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**Date of authorisation status change:**

20/06/1985

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

### Summary of Product Characteristics

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.

### Combined File of all Documents