

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:

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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:

•

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:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Romania

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:

8/01/2014

Manufacturing sites for batch release:

Orion Corporation

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

160156

Date of authorisation status change:

1/07/2024

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

Documents

Combined File of all Documents

Summary of Product Characteristics

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