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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
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:
Ireland

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:
12/03/2010

Manufacturing sites for batch release:
Orion Oyj / Orion Corporation / Orion Pharma

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10664/004/001

Date of authorisation status change:
12/03/2010

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