File downloaded on 2026-11-22
Source URL: https://medicines.health.europa.eu/veterinary/en/600000061186

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

Authorised

- DETOMIDINE HYDROCHLORIDE FOR VETERINARY USE

Product identification

Medicine name:
Domidine 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**
  - Milk. 12 hour
  - Meat and offal. 2 day
- **Horse**
  - Milk. 12 hour
  - Meat and offal. 2 day

Intravenous use:

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

Available in:
Netherlands

Package description:
(ID3) 20 millilitre(s): unspecified outer container with 1 Vial with 20 millilitre(s)
(ID2) 10 millilitre(s): unspecified outer container with 1 Vial with 10 millilitre(s)
(ID1) 5 millilitre(s): unspecified outer container with 1 Vial with 5 millilitre(s)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Eurovet Animal Health B.V.

Marketing authorisation date:
19/10/2011

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 10429

Date of authorisation status change:
7/02/2022

Reference member state:
Germany

Procedure number:
DE/V/0115/001

Concerned member states:
Austria Belgium Czechia Denmark France Hungary Ireland Italy Lithuania Luxembourg Netherlands Poland Portugal Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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

Summary of Product Characteristics

2400880-paren-20070123.pdf