

Medesedan 10 mg/ml Solution for Injection for Horses and Cattle

Authorised

- DETOMIDINE HYDROCHLORIDE FOR VETERINARY USE

Product identification

Medicine name:

Cepesedan RP 10 mg/ml, Solution for Injection for Horses and Cattle

Medesedan 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:

United Kingdom (Northern Ireland)

Available in:

United Kingdom (Northern Ireland)

Package description:

(ID4) 100 millilitre(s): unspecified outer container with 5 Vial (Glass) each with 20 millilitre(s), closed with Stopper and Cap (bromobutyl rubber`, Aluminium)

(ID3) 25 millilitre(s): unspecified outer container with 5 Vial (Glass) each with 5 millilitre(s), closed with Cap and Stopper (Aluminium, bromobutyl rubber`)

(ID2) 20 millilitre(s): unspecified outer container with 1 Vial (Glass) with 20 millilitre(s), closed with Stopper and Cap (bromobutyl rubber`, Aluminium)

(ID1) 5 millilitre(s): unspecified outer container with 1 Vial (Glass) with 5 millilitre(s), closed with Cap and Stopper (Aluminium, bromobutyl rubber`)

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:

CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:

3/08/2007

Manufacturing sites for batch release:

CP-Pharma Handelsgesellschaft mbH

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 20916/4002

Date of authorisation status change:

20/09/2023

Reference member state:

Germany

Procedure number:

DE/V/0117/001

Concerned member states:

Austria Belgium Czechia Denmark Finland France Hungary Ireland Italy
Latvia Lithuania Netherlands Norway Poland Portugal Slovakia Spain
Sweden United Kingdom (Northern Ireland)

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