SEDAXYLAN

Authorised

- Xylazine hydrochloride

Product identification

Medicine name:
SEDAXYLAN
Sedaxylan 20 mg/ml Oplossing voor injectie
Sedaxylan 20 mg/ml Solution injectable
Sedaxylan 20 mg/ml Injektionslösung

Active substance:
Xylazine hydrochloride

Target species:
Cattle
Horse
Dog
Cat

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Xylazine hydrochloride

23.30 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Milk. no withdrawal period zero days
  - Meat and offal. 1 day
- **Horse**
  - Meat and offal. 1 day
- **Dog**
- **Cat**

Intravenous use:

- **Cattle**
  - Milk. no withdrawal period zero days
  - Meat and offal. 1 day
- **Horse**
  - Meat and offal. 1 day
- **Dog**
- **Cat**

Subcutaneous use:

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Cattle

- Milk. no withdrawal period zero days
- Meat and offal. 1 day

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Horse

- Meat and offal. 1 day

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Dog

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Cat

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN05CM92

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Vial of 50 ml volume contents 50 ml. Glass type II, amber coloured. Bromobutyl rubber stopper type I secured with aluminium cap.
Vial of 30 ml volume contents 25 ml. Glass type II, amber coloured. Bromobutyl rubber stopper type I secured with aluminium cap.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Bibliographic application (Article 22 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Dechra Regulatory B.V.

Marketing authorisation date:
25/08/2003

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

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V254834

Date of authorisation status change:
25/08/2003

Reference member state:
Netherlands

Procedure number:
NL/V/0106/001

Concerned member states:
Austria Belgium Denmark France Germany Greece Ireland Italy Luxembourg Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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.
