SEDAXYLAN

Authorised

- Xylazine hydrochloride

Product identification

Medicine name:
SEDAXYLAN
Sedaxylan 20 mg/ml Injektionslösung für Hunde, Katzen, Pferde und Rinder

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

Intravenous use:

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

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05CM92

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

Authorisation status:
Valid

Authorised in:
Germany

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:
Eurovet Animal Health B.V.

Marketing authorisation date:
15/04/2003

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

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
400650.00.00

Date of authorisation status change:
5/02/2008

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

Combined File of all Documents

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