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# SEDAXYLAN

Authorised

- Xylazine hydrochloride

## Product identification

**Medicine name:**

SEDAXYLAN

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**Active substance:**

Xylazine hydrochloride

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**Target species:**

Cattle

Horse

Dog

Cat

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**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

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## Product details

**Active substance and strength:**

Xylazine hydrochloride

23.30 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intramuscular 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

**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

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**Horse**

- Meat and offal. 1 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN05CM92

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Belgium

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**Available in:**

Belgium

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

25/08/2003

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**Manufacturing sites for batch release:**

Eurovet Animal Health B.V.

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**Responsible authority:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

BE-V254834

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**Date of authorisation status change:**

25/08/2003

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0106/001

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**Concerned member states:**

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

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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.