

# Xylexx 20 mg/ml solution for injection for cattle, horses, dogs and cats

Authorised

- Xylazine

## Product identification

### Medicine name:

Xylexx 20 mg/ml solution for injection for cattle, horses, dogs and cats

Xylexx 20 mg/ml raztopina za injiciranje za govedo, konje, pse in mačke

### Active substance:

Xylazine

### Target species:

Cat

Cattle

Dog

Horse

### Route of administration:

Subcutaneous use

Intramuscular use

Intravenous use

## Product details

### Active substance and strength:

Xylazine

20.00 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

- Meat and offal. 1 day 1 day

#### Intravenous use:

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##### Horse

- Meat and offal. 1 day 1 day

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##### Cattle

- Meat and offal. 1 day 1 day

- Milk. no withdrawal period zero hours

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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:**

Slovenia

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**Package description:**

Cardboard box containing 1 vial of 30 ml

Cardboard box containing 5 vials of 30 ml

Polystyrene box containing 24 vials of 30 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Alfasan Nederland B.V.

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**Marketing authorisation date:**

24/10/2022

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**Manufacturing sites for batch release:**

Alfasan Nederland B.V.

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**Responsible authority:**

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

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**Authorisation number:**

DC/V/0760/001

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**Date of authorisation status change:**

24/10/2022

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0366/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania  
Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain  
Sweden 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.