

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

Authorised

- Xylazine hydrochloride

## Product identification

### Medicine name:

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

Xyla, 20 mg/ml süstelahus veistele, hobustele, koertele ja kassidele

### Active substance:

Xylazine hydrochloride

### Target species:

Cattle

Horse

Dog

Cat

### Route of administration:

Intravenous use

Intramuscular use

Subcutaneous use

## Product details

### Active substance and strength:

Xylazine hydrochloride  
23.32 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:**

**Intravenous use:**

• **Cattle**

- Meat and offal. 1 day
- Milk. 0 hour

• **Horse**

- Meat and offal. 1 day
- Milk. 0 hour

• **Dog**

**Intramuscular use:**

• **Cattle**

- Meat and offal. 1 day
- Milk. 0 hour

• **Dog**

• **Cat**

**Subcutaneous use:**

• **Cat**

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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:**

Estonia

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**Package description:**

50 ml clear, type II glass bottle, closed with a bromobutyl rubber stopper and secured with an aluminium cap. Package size: 5x50 ml in a cardboard box.

50 ml clear, type II glass bottle, closed with a bromobutyl rubber stopper and secured with an aluminium cap. Package size: 1x50 ml in a cardboard box.

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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:**

Interchemie Werken De Adelaar Eesti AS

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**Marketing authorisation date:**

24/09/2019

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**Manufacturing sites for batch release:**

Interchemie Werken De Adelaar Eesti AS

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**Responsible authority:**

State Agency Of Medicines

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**Authorisation number:**

2205

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**Date of authorisation status change:**

24/09/2019

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**Reference member state:**

Estonia

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**Procedure number:**

EE/V/0105/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Finland France

Germany Greece Hungary Iceland Ireland Italy Latvia Malta Netherlands  
Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden

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To consult adverse reactions on veterinary medicinal products please go to  
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## Documents

Summary of Product Characteristics

English (PDF)

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