Source URL: https://medicines.health.europa.eu/veterinary/en/600000027457

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 Sedachem 20,00 mg/ml Roztwór do wstrzykiwań

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)

Pharmaceutical form:

Solution for injection

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

Intramuscular use:

•

Cattle

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05CM92

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:
Poland
Package description:
Available only in <u>Polish</u>
Available only in <u>Polish</u>
Additional information
Entitlement type: Marketing Authorisation
Legal basis of product authorisation: Generic application (Article 13(1) of Directive No 2001/82/EC)
Marketing authorisation holder: Interchemie Werken De Adelaar Eesti AS
Marketing authorisation date: 25/01/2021
Manufacturing sites for batch release: Interchemie Werken De Adelaar Eesti AS
Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products
Authorisation number: 3066
Date of authorisation status change: 25/01/2021

Reference member state:

Estonia

Procedure number:

EE/V/0105/001

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

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 31/07/2025

Download

Package Leaflet

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