Xylased, 500mg, Lyofilizát a rozpouštědlo pro injekční roztok

Authorised

- Xylazine

Product identification

Medicine name:
Xylased, 500mg, Lyofilizát a rozpouštědlo pro injekční roztok

Active substance:
Xylazine

Target species:
Horse
Cattle
Fallow deer
Roe deer
Red deer

Route of administration:
Intravenous use
Intramuscular use

Product details

Active substance and strength:
Xylazine
500.00 milligram(s) / 1.00 Bottle

Pharmaceutical form:
Lyophilisate and solvent for solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Horse**
  - Meat and offal. no withdrawal period

Nepoužívat u koní, jejichž maso a mléko je určeno pro lidskou spotřebu.,

  - Milk. no withdrawal period

Nepoužívat u koní, jejichž maso a mléko je určeno pro lidskou spotřebu.,

- **Cattle**
  - Meat and offal. 3 day
  - Milk. 36 hour

Intramuscular use:

- **Horse**
  - Meat and offal. no withdrawal period

Nepoužívat u koní, jejichž maso a mléko je určeno pro lidskou spotřebu.,

  - Milk. no withdrawal period

Nepoužívat u koní, jejichž maso a mléko je určeno pro lidskou spotřebu.,

- **Cattle**
  - Meat and offal. 3 day
  - Milk. 36 hour

- **Fallow deer**

- Meat and offal. no withdrawal period

Nepoužívat u zvířat, jejichž maso je určeno pro lidskou spotřebu.,

- **Roe deer**
  - Meat and offal. no withdrawal period

Nepoužívat u zvířat, jejichž maso je určeno pro lidskou spotřebu.,

- **Red deer**
  - Meat and offal. no withdrawal period

Nepoužívat u zvířat, jejichž maso je určeno pro lidskou spotřebu.,

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Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN05CM92

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

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Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bioveta a.s.

Marketing authorisation date:
13/05/2010

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/032/10-C

Date of authorisation status change:
13/07/2021

To consult adverse reactions on veterinary medicinal products please go to 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.
