XYLAPAN, 20mg/ml, Injekční roztok

Authorised

- Xylazine

Product identification

Medicine name:
XYLAPAN, 20mg/ml, Injekční roztok

Active substance:
Xylazine

Target species:
Horse
Dog
Cattle
Cat

Route of administration:
Intravenous use
Intramuscular use

Product details

Active substance and strength:
Xylazine
20.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Horse**
  - Meat and offal. 1 day
  - Milk. 0 day
- **Dog**

Intramuscular use:

- **Cattle**
  - Meat and offal. 1 day
  - Milk. 0 day
- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05CM92

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Package description:

Available only in Czech

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetoquinol s.r.o.

Marketing authorisation date:
28/03/2001

Manufacturing sites for batch release:
Vetoquinol Biowet Sp. z o.o.

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

Authorisation number:
96/010/01-C

Date of authorisation status change:
23/08/2011

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

Documents

Summary of Product Characteristics

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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.
