XYLAZIN Ecuphar, 20mg/ml, Injekční roztok

Authorised

- Xylazine hydrochloride

Product identification

Medicine name:
XYLAZIN Ecuphar, 20mg/ml, Injekční roztok

Active substance:
Xylazine hydrochloride

Target species:
Horse
Cattle
Dog
Cat

Route of administration:
Intravenous use
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Xylazine hydrochloride

23.30 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 hour
- **Cattle**
  - 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:
Czechia

Available in:
Czechia

Package description:
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:
Ecuphar

Marketing authorisation date:
14/01/2002

Manufacturing sites for batch release:
Bioveta a.s.

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

Authorisation number:
96/006/02-C

Date of authorisation status change:
14/01/2002

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.