

XYLARIEM 20 mg/ml injekčný roztok

Authorised

- Xylazine hydrochloride

Product identification

Medicine name:

XYLARIEM 20 mg/ml injekčný roztok

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

23.30 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intravenous use:****• Cattle**

- Meat and offal. 0 day
Zero days

- Milk. 0 day
Zero days

• Horse

- Meat and offal. 0 day
Zero days

- Milk. 0 day
Zero days

• Dog**• Cat****Intramuscular use:****• Cattle**

- Meat and offal. 0 day
Zero days

- Milk. 0 day
Zero days

• Horse

- Meat and offal. 0 day
Zero days

- Milk. 0 day
Zero days

• Dog**• Cat****Subcutaneous use:****• Cattle**

- Meat and offal. 0 day
Zero days

- Milk. 0 day
Zero days

• Horse

- Meat and offal. 0 day
Zero days

- Milk. 0 day
Zero days

- **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:

Slovakia

Package description:

Available only in Slovak

Available only in Slovak

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Ecuphar

Marketing authorisation date:

26/04/2004

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Authorisation number:

96/057/04-S

Date of authorisation status change:

26/04/2004

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

Documents

Combined File of all Documents

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