

Xylasol 20 mg/ml, solution for injection for cattle, horses, dogs and cats

Authorised

- Xylazine hydrochloride

Product identification

Medicine name:

Xylasol 20 mg/ml, solution for injection for cattle, horses, dogs and cats
Xylavet 20 mg/ml Injektionslösung

Active substance:

Xylazine hydrochloride

Target species:

Cattle
Horse
Dog
Cat

Route of administration:

Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Xylazine hydrochloride

23.31 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Milk. no withdrawal period Should be 0 days
- Meat and offal. 1 day

-

Horse

- Milk. no withdrawal period Should be 0 days
- Meat and offal. 1 day

Intravenous use:

-

Cattle

- Milk. no withdrawal period Should be 0 days
- Meat and offal. 1 day

-

Horse

- Milk. no withdrawal period Should be 0 days
- Meat and offal. 1 day

Subcutaneous use:

-

Cattle

- Milk. no withdrawal period Should be 0 days
- Meat and offal. 1 day

-

Horse

- Milk. no withdrawal period Should be 0 days
- Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05CM92

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Available in:

Germany

Package description:

Cardboard box containing 1 vial (uncoloured glass type II, closed with bromobutyl rubber stopper, secured with aluminium cap) filled with 50 ml

Cardboard box containing 1 vial (uncoloured glass type II, closed with bromobutyl rubber stopper, secured with aluminium cap) filled with 25 ml

Cardboard box containing 1 vial (uncoloured glass type II, closed with bromobutyl rubber stopper, secured with aluminium cap) filled with 10 ml

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:

CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:

19/04/2012

Manufacturing sites for batch release:

CP-Pharma Handelsgesellschaft mbH

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

401510.00.00

Date of authorisation status change:

27/04/2018

Reference member state:

Netherlands

Procedure number:

NL/V/0158/001

Concerned member states:

Austria Denmark Finland France Germany Hungary Norway Spain Sweden

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

Documents

Combined File of all Documents

Summary of Product Characteristics

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108964 - par.pdf