

XYLAPAN 20 mg/mL otopina za injekciju, za konja, govedo, psa i mačku

Not
authorised

- Xylazine hydrochloride

Product identification

Medicine name:

XYLAPAN 20 mg/mL otopina za injekciju, za konja, govedo, psa i mačku

Active substance:

Xylazine hydrochloride

Target species:

Cattle

Dog

Cat

Horse

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Xylazine hydrochloride
20.00 milligram(s)/millilitre / 1.00 Vial

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. 1 day
- Milk. no withdrawal period N/A

Intravenous use:

-

Horse

- Meat and offal. 3 day
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05CM92

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Croatia

Package description:

Available only in Croatian

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:

Vetoquinol S.A.

Marketing authorisation date:

3/07/2015

Manufacturing sites for batch release:

Vetoquinol Biowet Sp. z o.o.

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/13-01/600

Date of authorisation status change:

31/08/2023

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.