File downloaded on 2026-01-03
Source URL: https://medicines.health.europa.eu/veterinary/en/600000107582

XYLAPAN, 20 mg/ml ενέσιμο διάλυμα για ιπποειδή, βοοειδή, σκύλους και γάτες

Authorised

- Xylazine hydrochloride

Product identification

Medicine name:
XYLAPAN, 20 mg/ml ενέσιμο διάλυμα για ιπποειδή, βοοειδή, σκύλους και γάτες

Active substance:
Xylazine hydrochloride

Target species:
Horse
Cattle
Dog
Cat

Route of administration:
Intravenous use
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Xylazine hydrochloride
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. 3 day

Intramuscular use:

- **Cattle**
  - Meat and offal, milk. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN05CM92

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

Available in:
Greece

Package description:
Available only in Greek

Available only in Greek

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
Vetoquinol S.A.

Marketing authorisation date:
7/05/1997

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

Responsible authority:
National Organization For Medicines

Authorisation number:
51834/13-07-2012/K-0114001

Date of authorisation status change:
5/05/2020

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