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SEDAXYLAN

Authorised

- Xylazine hydrochloride

Product identification

Medicine name:

SEDAXYLAN

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

23.30 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Cattle

- Milk. no withdrawal period zero days

- Meat and offal. 1 day

-

Horse

- Meat and offal. 1 day

Intravenous use:

-

Cattle

- Milk. no withdrawal period zero days

- Meat and offal. 1 day

-

Horse

- Meat and offal. 1 day

Subcutaneous use:

-

Cattle

- Milk. no withdrawal period zero days

- Meat and offal. 1 day

-

Horse

- 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:

Italy

Available in:

Italy

Package description:

Vial of 50 ml volume contents 50 ml. Glass type II, amber coloured. Bromobutyl rubber stopper type I secured with aluminium cap.

Vial of 30 ml volume contents 25 ml. Glass type II, amber coloured. Bromobutyl rubber stopper type I secured with aluminium cap.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Bibliographic application (Article 22 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Eurovet Animal Health B.V.

Marketing authorisation date:

26/02/2004

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Ministry Of Health

Authorisation number:

103595

Date of authorisation status change:

26/02/2004

Reference member state:

Netherlands

Procedure number:

NL/V/0106/001

Concerned member states:

Austria Belgium Denmark France Germany Greece Ireland Italy
Luxembourg Portugal Spain United Kingdom (Northern Ireland)

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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