

File downloaded on 2026-04-22

Source URL: <https://medicines.health.europa.eu/veterinary/en/600000037662>

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:

France

Available in:

France

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:

23/05/2003

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/1427734 4/2003

Date of authorisation status change:

23/05/2008

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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.