

# Nerfasin vet. 100 mg/ml, solution for injection for cattle and horses

Authorised

- Xylazine hydrochloride

## Product identification

**Medicine name:**

Nerfasin vet. 100 mg/ml, solution for injection for cattle and horses

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**Active substance:**

Xylazine hydrochloride

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**Target species:**

Cattle

Horse

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**Route of administration:**

Intramuscular use

Intravenous use

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## Product details

**Active substance and strength:**

Xylazine hydrochloride

116.55 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Cattle**

- Milk. no withdrawal period zero days

- Meat and offal. 1 day

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**Horse**

- Milk. no withdrawal period zero days

- Meat and offal. 1 day

**Intravenous use:**

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**Cattle**

- Milk. no withdrawal period zero days

- Meat and offal. 1 day

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**Horse**

- Milk. no withdrawal period zero days

- Meat and offal. 1 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN05CM92

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Luxembourg

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**Package description:**

50 ml product in a 50 ml clear type II glass vials closed with a bromobutyl rubber stopper and aluminium cap in a carton box.

25 ml product in a 30 ml clear type II glass vials closed with a bromobutyl rubber stopper and aluminium cap in a carton box.

10 ml product in a 10 ml clear type II glass vials closed with a bromobutyl rubber stopper and aluminium cap in a carton box.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

22/03/2013

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

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**Responsible authority:**

Ministry Of Health And Social Security

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**Authorisation number:**

V 914/13/05/1304

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**Date of authorisation status change:**

22/03/2013

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0157/002

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**Concerned member states:**

Austria Belgium Czechia Denmark Finland France Greece Hungary Iceland  
Ireland Italy Luxembourg Norway Poland Portugal Slovakia Spain Sweden  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

108959 - par.pdf