

# TILDREN 5 MG/ML POWDER AND SOLVENT FOR SOLUTION FOR INJECTION FOR HORSES

Authorised

- Tiludronate disodium hemihydrate

## Product identification

**Medicine name:**

TILDREN 5 MG/ML POWDER AND SOLVENT FOR SOLUTION FOR INJECTION FOR HORSES

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**Active substance:**

Tiludronate disodium hemihydrate

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**Target species:**

Horse

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**Route of administration:**

Intravenous use

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## Product details

**Active substance and strength:**

Tiludronate disodium hemihydrate  
5.83 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Powder and solvent for solution for injection

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

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**Horse**

- Meat and offal. 0 day
- Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

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

QM05BA05

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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:**

Netherlands

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**Available in:**

Netherlands

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**Package description:**

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Audevard

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**Marketing authorisation date:**

4/06/2003

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**Manufacturing sites for batch release:**

Ceva Sante Animale

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 10079

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**Date of authorisation status change:**

21/03/2022

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**Reference member state:**

France

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**Procedure number:**

FR/V/0134/001

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**Concerned member states:**

Austria Greece Italy Luxembourg Netherlands Norway Spain

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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

English (PDF)

Published on: 14/03/2026

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