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
TILDREN®

Active substance:
Tiludronate disodium hemihydrate

Target species:
Horse

Route of administration:
Intravenous use

Product details

Active substance and strength:
Tiludronate disodium hemihydrate
5.83 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Powder and solvent for solution for injection

Withdrawal period by route of administration:

Intravenous use:

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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM05BA05

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

Authorisation status:
Valid

Authorised in:
Norway

Package description:
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Audevard

Marketing authorisation date:
1/08/2003

Manufacturing sites for batch release:
CEVA Santé Animale

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
02-1588

Date of authorisation status change:
9/01/2007

Reference member state:
France

Procedure number:
FR/V/0134/001

Concerned member states:
Austria Greece Italy Luxembourg Netherlands Norway Spain

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

Documents

Summary of Product Characteristics

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Package Leaflet

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