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Osphos 51 mg/ml solution for injection for horses

Authorised

- Clodronate disodium tetrahydrate

Product identification

Medicine name:
Osphos 51 mg/ml solution for injection for horses

Active substance:
Clodronate disodium tetrahydrate

Target species:
Horse

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Clodronate disodium tetrahydrate
74.98 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Horse**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM05BA02

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

Authorisation status:
Valid

Authorised in:
Sweden

Available in:
Sweden

Package description:
Clear glass (type I) vial with siliconized rubber stopper, an aluminium seal and a plastic flip-off cap containing 15 ml of clodronic acid solution.Each cardboard box contains 1 vial

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Dechra Regulatory B.V.

Marketing authorisation date:

24/09/2015

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
51366

Date of authorisation status change:
24/09/2015

Reference member state:
Ireland

Procedure number:
IE/V/0508/001

Concerned member states:
Austria Belgium Czechia Finland France Germany Hungary Iceland Italy Luxembourg Netherlands Poland Portugal Slovakia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 26/01/2025
Updated on: 14/03/2026

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Package Leaflet

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Combined File of all Documents

Labelling

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