

PHOSPHONORTONIC SOLUTION INJECTABLE

Authorised

- Toldimfos sodium trihydrate

Product identification

Medicine name:

PHOSPHONORTONIC SOLUTION INJECTABLE

Active substance:

Toldimfos sodium trihydrate

Target species:

Cattle

Pig

Cat

Horse

Horse (mare)

Sheep

Goat

Dog

Route of administration:

Intramuscular use

Subcutaneous use

Intravenous use

Product details

Active substance and strength:

Toldimfos sodium trihydrate

180.29 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Milk. 0 day

- Meat and offal. 0 day

-

Pig

- Meat and offal. 0 day

-

Horse

- Meat and offal. 0 day

-

Horse (mare)

- Milk. 0 day

-

Sheep

- Milk. 0 day

- Meat and offal. 0 day

-

Goat

- Milk. 0 day

- Meat and offal. 0 day

Subcutaneous use:

•

Cattle

- Milk. 0 day

- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

•

Horse

- Meat and offal. 0 day

•

Horse (mare)

- Milk. 0 day

•

Sheep

- Milk. 0 day

- Meat and offal. 0 day

•

Goat

- Milk. 0 day

- Meat and offal. 0 day

Intravenous use:

•

Cattle

- Milk. 0 day

- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

•

Horse

- Meat and offal. 0 day

•

Horse (mare)

- Milk. 0 day

•

Sheep

- Milk. 0 day

- Meat and offal. 0 day

•

Goat

- Milk. 0 day

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12CX90

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

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:

Vetoquinol S.A.

Marketing authorisation date:

19/06/1992

Manufacturing sites for batch release:

Vetoquinol S.A.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/6904755 8/1992

Date of authorisation status change:

19/06/2012

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.