

COFACALCIUM SOLUTION INJECTABLE

Authorised

- Calcium gluconate monohydrate
- Magnesium hypophosphite hexahydrate

Product identification

Medicine name:

COFACALCIUM SOLUTION INJECTABLE

Active substance:

Calcium gluconate monohydrate

Magnesium hypophosphite hexahydrate

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:

Calcium gluconate monohydrate
280.00 milligram(s) / 1.00 millilitre(s)

Magnesium hypophosphite hexahydrate
90.00 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:

QA12AX

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

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:

Dopharma France S.A.S.

Marketing authorisation date:

21/05/1984

Manufacturing sites for batch release:

Dopharma France

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/5869837 4/1984

Date of authorisation status change:

21/05/2009

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.