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Cofacalcium oldatos injekció A.U.V.

Not authorised

- Boric acid
- Magnesium hypophosphite hexahydrate
- Calcium gluconate

Product identification

Medicine name:
Cofacalcium oldatos injekció A.U.V.

Active substance:
Boric acid

Magnesium hypophosphite hexahydrate

Calcium gluconate

Target species:
Cattle
Sheep
Goat
Pig
Horse
Dog
Cat

Route of administration:
Intramuscular use

Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Boric acid
5.75 gram(s) / 100.00 millilitre(s)

Magnesium hypophosphite hexahydrate
9.00 gram(s) / 100.00 millilitre(s)

Calcium gluconate
28.00 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
- **Goat**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day

Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
- **Goat**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
- **Goat**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
-

Horse

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA12AX

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

Authorisation status:
Surrendered

Authorised in:
Hungary

Package description:
Available only in Hungarian
Available only in Hungarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Dopharma Research B.V.

Marketing authorisation date:
30/05/1997

Manufacturing sites for batch release:
Dopharma France

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
12/08/2024

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

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