CALCIUMVIT B12

Authorised

- Cyanocobalamin
- Calcium gluconate
- Magnesium gluconate

Product identification

Medicine name:
CALCIUMVIT B12

Active substance:
Cyanocobalamin
Calcium gluconate
Magnesium gluconate

Target species:
Cattle
Dog
Cat
Pig
Horse

Route of administration:
Intraperitoneal use
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Cyanocobalamin
0.04 milligram(s) / 1.00 millilitre(s)

Calcium gluconate
200.00 milligram(s) / 1.00 millilitre(s)

Magnesium gluconate
60.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intraperitoneal use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day

Intramuscular use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
-

Pig

- Meat and offal. 0 day

•

Horse

- 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

Subcutaneous use:

•

Cattle

- Milk. 0 day

- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

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Horse

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11BA

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

Authorisation status:
Valid

Authorised in:
Italy

Available in:
Italy

Package description:
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian

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:
Fatro S.p.A.

Marketing authorisation date:
8/06/1962

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Ministry Of Health

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

Date of authorisation status change:
10/08/2001

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

Documents

Combined File of all Documents

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