VITAMINE B12 VETOQUINOL

Authorised

Cyanocobalamin

Product identification

Medicine name:

VITAMINE B12 VETOQUINOL

Active substance:

Cyanocobalamin

Target species:

Cattle Pig Cat Horse Horse (mare) Sheep Dog

Route of administration:

Intramuscular use Subcutaneous use Oral use Intravenous use

Product details

Active substance and strength:

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

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration: Intramuscular use:

Cattle

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

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Pig

- Meat and offal. 0 day
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Cat

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Horse

- Meat and offal. 0 day
- •

Horse (mare)

- Milk. 0 day

Sheep

- Milk. 0 day
- Meat and offal. 0 day
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Dog

Subcutaneous use:

• Cattle - Meat and offal. 0 day

- Milk. 0 day

Pig

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- Meat and offal. 0 day

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Cat

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Horse

- Meat and offal. 0 day

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Horse (mare)

- Milk. 0 day

Sheep

- Meat and offal. 0 day

- Milk. 0 day

•

Dog

Oral use:

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Cattle

- Meat and offal. 0 day

- Milk. 0 day

• Pig

- Meat and offal. 0 day

• Cat

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Horse

- Meat and offal. 0 day

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Horse (mare)

- Milk. 0 day

•

Sheep

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

•

Dog

Intravenous use:

• Cattle

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

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Pig

- Meat and offal. 0 day

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Cat

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Horse

- Meat and offal. 0 day

Horse (mare)

- Milk. 0 day

Sheep

•

- Meat and offal. 0 day

- Milk. 0 day

Dog

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Anatomical therapeutic chemical veterinary (ATCvet) codes: QB03BA01

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:

Vetoquinol S.A.

Marketing authorisation date:

19/06/1992

Manufacturing sites for batch release: VETOQUINOL

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number: FR/V/5403932 4/1992

Date of authorisation status change:

18/12/2023

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

Documents

Summary of Product Characteristics

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