

# BICARBOSSILASI, 40 mg/ml liofilizzato per soluzione iniettabile per bovini, equini e cani

Authorised

- THIAMINE-DIPHOSPHATE CHLORIDE

## Product identification

### Medicine name:

BICARBOSSILASI, 40 mg/ml liofilizzato per soluzione iniettabile per bovini, equini e cani

### Active substance:

THIAMINE-DIPHOSPHATE CHLORIDE

### Target species:

Cattle

Horse

Dog

### Route of administration:

Intramuscular use

Intravenous use

Subcutaneous use

## Product details

### Active substance and strength:

THIAMINE-DIPHOSPHATE CHLORIDE

400.00 milligram(s) / 10.00 millilitre(s)

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### Pharmaceutical form:

Lyophilisate and solvent for solution for injection

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### Withdrawal period by route of administration:

#### Intramuscular use:

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##### Cattle

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

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##### Cattle

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

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##### Horse

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

#### Intravenous use:

- 

##### Cattle

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

- 

##### Cattle

- Milk. 0 day

- Meat and offal. 0 day

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**Horse**

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

**Subcutaneous use:**

- 

**Cattle**

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

- 

**Cattle**

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

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**Horse**

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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QA11DA

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**Legal status of supply:**

Veterinary medicinal product not subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Italy

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**Package description:**

Available only in Italian

Available only in Italian

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Teknofarma S.r.l.

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**Marketing authorisation date:**

2/05/1960

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**Manufacturing sites for batch release:**

Teknofarma S.r.l.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

1/01/2009

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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