

Calcibel Forte, 380/60/50 mg/ml solution for infusion for horses, cattle, sheep, goats and pigs

Authorised

- Calcium gluconate
- Magnesium chloride hexahydrate
- Boric acid

Product identification

Medicine name:

Calcibel Forte, 380/60/50 mg/ml solution for infusion for horses, cattle, sheep, goats and pigs

Active substance:

Calcium gluconate

Magnesium chloride hexahydrate

Boric acid

Target species:

Horse

Cattle

Sheep

Goat

Pig

Route of administration:

Intravenous use

Product details

Active substance and strength:

Calcium gluconate

380.00 milligram(s) / 1.00 millilitre(s)

Magnesium chloride hexahydrate

60.00 milligram(s) / 1.00 millilitre(s)

Boric acid

50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for infusion

Withdrawal period by route of administration:

Intravenous use:

-

Horse

- Milk. 0 hour

- Meat and offal. 0 day

-

Cattle

- Milk. 0 hour

- Meat and offal. 0 day

-

Sheep

- Milk. 0 hour

- Meat and offal. 0 day

-

Goat

- Milk. 0 hour

- Meat and offal. 0 day

-

Pig

- 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:

Valid

Authorised in:

Greece

Available in:

Greece

Package description:

Plastic Bottle 12 x 500.0 millilitre(s)

Plastic Bottle 1 x 500.0 millilitre(s)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Bela-Pharm GmbH & Co. KG

Marketing authorisation date:

20/09/2021

Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

Responsible authority:

National Organization For Medicines

Authorisation number:

84808/21-09-2021/K-0214302

Date of authorisation status change:

20/09/2021

Reference member state:

Czechia

Procedure number:

CZ/V/0170/001

Concerned member states:

Austria Cyprus Denmark Estonia Finland Greece Hungary Iceland Ireland
Italy Latvia Lithuania Poland Portugal Romania Slovakia Slovenia Spain
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 14/03/2026

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Package Leaflet

English (PDF)

Published on: 14/03/2026

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Labelling