NOVACOC FORTE soluție perfuzabilă pentru cai, bovine, porci

Authorised

- Sodium dihydrogen phosphate dihydrate
- Metamizole sodium
- Magnesium gluconate
- Glucose
- Calcium gluconate
- Caffeine
- Acetylmethionine

Product identification

Medicine name:
NOVACOC FORTE soluție perfuzabilă pentru cai, bovine, porci

Active substance:
Sodium dihydrogen phosphate dihydrate

Metamizole sodium

Magnesium gluconate

Glucose

Calcium gluconate

Caffeine

Acetylmethionine

Target species:

Horse
Cattle
Pig

Route of administration:
Intravenous use

Product details

Active substance and strength:
Sodium dihydrogen phosphate dihydrate
4.02 milligram(s) / 1.00 millilitre(s)

Metamizole sodium
40.00 milligram(s) / 1.00 millilitre(s)

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

Glucose
181.82 milligram(s) / 1.00 millilitre(s)

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

Caffeine
3.50 milligram(s) / 1.00 millilitre(s)

Acetylmethionine
40.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for infusion

Withdrawal period by route of administration:
Intravenous use:

- **Horse**
  - Meat and offal. 6 day

Do not use in milk-producing mares for human consumption.

- **Cattle**
  - Meat and offal. 13 day
  - Milk. 3 day
- **Pig**
  - Meat and offal. 6 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02BB52

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

Authorisation status:
Valid

Authorised in:
Romania

Available in:
Romania

Package description:
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Vetviva Richter GmbH

Marketing authorisation date:
2/02/2004

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
230038

Date of authorisation status change:
17/12/2025

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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