Novacoc forte - Infusion Solution for Animals

Authorised

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

Product identification

Medicine name:
Новакок форте - разтвор за инфузия за животни
Novacoc forte - Infusion Solution for Animals

Active substance:
Metamizole sodium
Caffeine
Calcium gluconate
Magnesium gluconate
Sodium dihydrogen phosphate dihydrate
Acetylmethionine
Glucose monohydrate

Target species:
Horse
Cattle

Cattle (calf)
Pig

Route of administration:
Intravenous use

Product details

Active substance and strength:
Metamizole sodium
4.00 gram(s) / 100.00 millilitre(s)

Caffeine
0.35 gram(s) / 100.00 millilitre(s)

Calcium gluconate
10.00 gram(s) / 100.00 millilitre(s)

Magnesium gluconate
1.00 gram(s) / 100.00 millilitre(s)

Sodium dihydrogen phosphate dihydrate
0.40 gram(s) / 100.00 millilitre(s)

Acetylmethionine
4.00 gram(s) / 100.00 millilitre(s)

Glucose monohydrate
18.18 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Horse**
  - Meat and offal. 5 day
- **Cattle**
  - Meat and offal. 6 day
  - Milk. 3 day Мляко: 2 ½ дни (5 издоявания).

- **Cattle (calf)**
  - Meat and offal. 6 day
- **Pig**
  - Meat and offal. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02BB52

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetviva Richter GmbH

Marketing authorisation date:
This information is not available for this product.

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Bulgarian Agency For Food Safety

Authorisation number:
0022-2184

Date of authorisation status change:
4/12/2008

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

Documents

Summary of Product Characteristics

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Labelling

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

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