Novacoc Forte, otopina za infuziju, konj, govedo i svinja

Authorised

- Metamizole sodium
- Caffeine
- CALCIUM GLUCONATE FOR INJECTION
- Magnesium gluconate
- SODIUM DIHYDROGEN PHOSPHATE DIHYDRATE PH. EUR.
- Acetylmethionine
- Glucose monohydrate

Product identification

Medicine name:
Novacoc Forte, otopina za infuziju, konj, govedo i svinja

Active substance:
Metamizole sodium

Caffeine

CALCIUM GLUCONATE FOR INJECTION

Magnesium gluconate

SODIUM DIHYDROGEN PHOSPHATE DIHYDRATE PH. EUR.

Acetylmethionine

Glucose monohydrate

Target species:
Horse

Cattle
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 FOR INJECTION
10.00 gram(s) / 100.00 millilitre(s)

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

SODIUM DIHYDROGEN PHOSPHATE DIHYDRATE PH. EUR.
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 infusion

Withdrawal period by route of administration:
Intravenous use:

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

Ne smije se primjenjivati kobilama čije se mlijeko koristi za hranu.

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Cattle

- Meat and offal. 13 day
- Milk. 60 hour

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Pig

- Meat and offal. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02BB52

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

Authorisation status:
Valid

Authorised in:
Croatia

Available in:
Croatia

Package description:
Available only in Croatian

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Vetviva Richter GmbH

Marketing authorisation date:

4/06/2020

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/20-01/225

Date of authorisation status change:
11/08/2023

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

Documents

Summary of Product Characteristics

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