ANALGIN 500 mg/ml injekčný roztok

Not authorised

- Metamizole sodium

Product identification

Medicine name:
ANALGIN 500 mg/ml injekčný roztok

Active substance:
Metamizole sodium

Target species:
Cattle
Horse
Dog

Route of administration:
Intravenous use
Intramuscular use

Product details

Active substance and strength:
Metamizole sodium
500.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Cattle**
  - Meat and offal. 12 day
  - Milk. 96 hour 8 milkings
- **Horse**
  - All relevant tissues. no withdrawal period

Do not use in horses whose meat is intended for human consumption.

Intramuscular use:

- **Cattle**
  - Meat and offal. 12 day
  - Milk. 96 hour 8 milkings
- **Horse**
  - All relevant tissues. no withdrawal period

Do not use in horses whose meat is intended for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02BB02

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

Authorisation status:
Surrendered

Authorised in:
Slovakia

Package description:
Available only in Slovak
Available only in Slovak
Available only in Slovak

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
BB Pharma a.s.

Marketing authorisation date:
1/08/1996

Manufacturing sites for batch release:
Farmacia Martin a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
07/795/69-S

Date of authorisation status change:
11/11/2022

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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