Buscopan Compositum ad us.vet. 4 mg/ml + 500 mg/ml, solution injectable pour chevaux, veaux et chiens

Authorised

- Hyoscine butylbromide
- Metamizole sodium

Product identification

Medicine name:
Buscopan Compositum ad us.vet. 4 mg/ml + 500 mg/ml, solution injectable pour chevaux, veaux et chiens
Buscopan Compositum ad us. vet. 4 mg/ml + 500 mg/ml, Injektionslösung für Pferde, Kälber und Hunde

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Active substance:
Hyoscine butylbromide
Metamizole sodium

Target species:
Cattle
Cattle (calf)
Horse
Dog

Route of administration:

Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Hyoscine butylbromide
4.00 milligram(s) / 1.00 millilitre(s)

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Cattle**
  - Milk. no withdrawal period

Not to be used in animals producing milk for human consumption

- **Cattle (calf)**
  - Meat and offal. 15 day

- **Horse**
  - Meat and offal. 12 day
  - Milk. no withdrawal period

Not to be used in animals producing milk for human consumption

Subcutaneous use:

- **Cattle**

- Milk. no withdrawal period

Not to be used in animals producing milk for human consumption

- **Cattle (calf)**
  - Meat and offal. 15 day

- **Horse**
  - Meat and offal. 12 day
  - Milk. no withdrawal period

Not to be used in animals producing milk for human consumption

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA03BB01

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

Authorisation status:
Valid

Authorised in:
Luxembourg

Available in:
Luxembourg

Package description:
Buscopan Compositum ad us. vet 1 Vial of 100 ml with Solution for injection

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:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
31/12/1991

Manufacturing sites for batch release:
LABIANA LIFE SCIENCES, S.A.

Responsible authority:
Ministry Of Health And Social Security

Authorisation number:
V 642/97/11/0345

Date of authorisation status change:
16/03/2010

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

Documents

Summary of Product Characteristics

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Package Leaflet

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