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Adrenaline, 1 mg/ml oplossing voor injectie

Not authorised

- Adrenaline

Product identification

Medicine name:
Adrenaline, 1 mg/ml oplossing voor injectie

Active substance:
Adrenaline

Target species:
Cattle
Sheep
Horse
Pig
Dog

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Adrenaline
1.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. no withdrawal period
  - Meat and offal. 1 day
- **Sheep**
  - Milk. no withdrawal period
  - Meat and offal. 1 day
- **Horse**
  - Meat and offal. 1 day
- **Pig**
  - Meat and offal. 1 day

Intravenous use:

- **Cattle**
  - Milk. no withdrawal period
  - Meat and offal. 1 day
- **Horse**
  - Meat and offal. 1 day

Subcutaneous use:

- **Cattle**
  - Milk. no withdrawal period
  - Meat and offal. 1 day
- **Sheep**
  - Milk. no withdrawal period
  - Meat and offal. 1 day
- **Horse**
  - Meat and offal. 1 day
- **Pig**
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC01CA24

Legal status of supply:
Medicinal product subject to medical prescription

Authorisation status:
Surrendered

Authorised in:
Netherlands

Package description:
Available only in Dutch
Available only in Dutch

Additional information

Entitlement type:

Marketing Authorisation

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

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
8/07/1992

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 4756

Date of authorisation status change:
18/04/2024

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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