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Adrilan 1 mg/ml solução injectável

Authorised

- Adrenaline

Product identification

Medicine name:
Adrilan 1 mg/ml solução injectável

Active substance:
Adrenaline

Target species:
Dog
Horse
Cattle
Sheep
Cat
Ornamental bird

Route of administration:
Intramuscular use
Intravenous use
Intracardiac 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:

- **Dog**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day

Intravenous use:

- **Dog**
  - Meat and offal. 0 day
- **Cat**

- Meat and offal. 0 day

- **Horse**
  - Meat and offal. 0 day

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day

- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day

- **Ornamental bird**
  - Meat and offal. 0 day

Não é autorizada a administração a aves produtoras de ovos para consumo humano.

Intracardiac use:

- **Dog**
  - Meat and offal. 0 day

- **Cat**
  - Meat and offal. 0 day

- **Ornamental bird**
  - Meat and offal. 0 day

Não é autorizada a administração a aves produtoras de ovos para consumo humano.

Subcutaneous use:

-

Dog

- Meat and offal. 0 day

•

Cat

- Meat and offal. 0 day

•

Cattle

- Meat and offal. 0 day
- Milk. 0 day

•

Horse

- Meat and offal. 0 day

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Sheep

- Meat and offal. 0 day
- Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QR03AA01

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

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Authorisation status:
Valid

Authorised in:
Portugal

Package description:
Available only in Portuguese

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:
Medinfar-Serological Products E Equipamentos S.A.

Marketing authorisation date:
12/05/2000

Manufacturing sites for batch release:
Labesfal Laboratorios Almiro S.A.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
51322

Date of authorisation status change:
1/07/2010

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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