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Adrenaline, 1 mg/ml oplossing voor injectie

Not authorised

• Epinephrine

Product identification

Medicine name:

Adrenaline, 1 mg/ml oplossing voor injectie

Active substance:

Epinephrine

Target species:

Cattle

Sheep

Horse

Pig

Dog

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

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

•

Dog

Intravenous use:

•

Cattle

- Milk. no withdrawal period
- Meat and offal. 1 day

•

Sheep

•

Horse - Meat and offal. 1 day Pig Dog Subcutaneous use: Cattle - Milk. no withdrawal period - Meat and offal. 1 day Sheep - Milk. no withdrawal period - Meat and offal. 1 day Meat and offal. 1 day

Horse

- Meat and offal. 1 day

•

Pig

- Meat and offal. 1 day

•

Dog

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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