

Dexatat 2 mg/ml solution for Injection

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:

Dexatat 2 mg/ml solution for Injection

Active substance:

Dexamethasone sodium phosphate

Target species:

Cattle

Dog

Horse

Cat

Pig

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Dexamethasone sodium phosphate

2.63 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

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Cattle

- Meat and offal. 16 day

- Milk. 84 hour

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Horse

- Meat and offal. 6 month

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Pig

- Meat and offal. 4 day

Intravenous use:

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Cattle

- Meat and offal. 16 day

- Milk. 84 hour

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Pig

- Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

Vial: 100 ml amber glass vial.Multipack: 12 x 100 ml.Stopper: Rubber.Cap: Aluminium Flanging cap.

Vial: 50 ml amber glass vial.Multipack: 12 x 50 ml.Stopper: Rubber.Cap: Aluminium Flanging cap.

Vial: 100 ml amber glass vial.Stopper: Rubber.Cap: Aluminium Flanging cap.

Vial: 50 ml amber glass vial.Stopper: Rubber.Cap: Aluminium Flanging cap.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

aniMedica GmbH

Marketing authorisation date:

1/10/1989

Manufacturing sites for batch release:

aniMedica Herstellungs GmbH

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10826/002/001

Date of authorisation status change:

1/10/1989

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

Documents

Summary of Product Characteristics