

Dexamethason ad us.vet. 2 mg/ml Injektionslösung für Pferde, Rinder, Schweine, Hunde und Katzen

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:

Dexamethason ad us.vet. 2 mg/ml Injektionslösung für Pferde, Rinder, Schweine, Hunde und Katzen

Active substance:

Dexamethasone sodium phosphate

Target species:

Cattle

Dog

Horse

Cat

Pig

Route of administration:

Intravenous use

Intraarticular use

Intramuscular 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:

Intravenous use:

-

Cattle

- Meat and offal. 16 day

- Milk. 4 day

-

Horse

- Meat and offal. 16 day

- Milk. no withdrawal period

Stuten, von denen Milch als Lebensmittel gewonnen werden soll, sind von der Anwendung auszuschliessen.

-

Pig

- Meat and offal. 4 day

Intraarticular use:

-

Cattle

- Meat and offal. 16 day

- Milk. 4 day

-

Horse

- Meat and offal. 16 day
- Milk. no withdrawal period

Stuten, von denen Milch als Lebensmittel gewonnen werden soll, sind von der Anwendung auszuschliessen.

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Pig

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Intramuscular use:

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Horse

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Stuten, von denen Milch als Lebensmittel gewonnen werden soll, sind von der Anwendung auszuschliessen.

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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:

Germany

Available in:

Germany

Package description:

Available only in [German](#)

Available only in [German](#)

Available only in [German](#)

Available only in [German](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:

CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:

25/02/2005

Manufacturing sites for batch release:

Produlab Pharma B.V.

CP-Pharma Handelsgesellschaft mbH

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

6684978.00.00

Date of authorisation status change:

25/02/2005

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

Documents

Combined File of all Documents

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