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.

- **Pig**
  - Meat and offal. 4 day

Intramuscular 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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

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

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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.