

Dexamethason 4 mg/ml

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:

Dexamethason 4 mg/ml

Active substance:

Dexamethasone sodium phosphate

Target species:

Dog
Cat
Horse
Cattle
Pig

Route of administration:

Intramuscular use
Subcutaneous use
Intravenous use

Product details

Active substance and strength:

Dexamethasone sodium phosphate
5.24 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Horse

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

Nicht bei Stuten anwenden, deren Milch für den menschlichen Verzehr vorgesehen ist.

-

Cattle

- Meat and offal. 16 day
- Milk. 4 day

-

Pig

- Meat and offal. 4 day

Subcutaneous use:

-

Cattle

- Meat and offal. 16 day
- Milk. 4 day

-

Horse

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

Nicht bei Stuten anwenden, deren Milch für den menschlichen Verzehr vorgesehen ist.

-

Pig

- Meat and offal. 4 day

Intravenous use:

-

Horse

- Meat and offal. 16 day

- Milk. no withdrawal period

Nicht bei Stuten anwenden, deren Milch für den menschlichen Verzehr vorgesehen ist.

-

Cattle

- Meat and offal. 16 day

- Milk. 4 day

-

Pig

- Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA16QA

QD07AB

QH02AB02

QM09AX

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](#)

Available only in [German](#)

Available only in [German](#)

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Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:

Bela-Pharm GmbH & Co. KG

Marketing authorisation date:

19/12/2005

Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

6933074.00.00

Date of authorisation status change:

19/12/2005

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.