Dexadreson forte 1,32 mg/ml + 2,67 mg/ml Injektionssuspension für Rinder, Pferde und Hunde

Authorised

- Dexamethasone sodium phosphate
- DEXAMETHASONE 21-PHENYLPROPIONATE

Product identification

Medicine name:
Dexadreson forte 1,32 mg/ml + 2,67 mg/ml Injektionssuspension für Rinder, Pferde und Hunde

Active substance:
Dexamethasone sodium phosphate
DEXAMETHASONE 21-PHENYLPROPIONATE

Target species:
Cattle
Dog
Horse

Route of administration:
Periarticular use
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Dexamethasone sodium phosphate
1.32 milligram(s) / 1.00 millilitre(s)

DEXAMETHASONE 21-PHENYLPROPIONATE
2.67 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Periarticular use:

- **Cattle**
  - Milk. 7 day
  - Meat and offal. 48 day
- **Horse**
  - Milk. no withdrawal period

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

  - Meat and offal. 47 day

Subcutaneous use:

- **Cattle**
  - Milk. 7 day
  - Meat and offal. 48 day
- **Horse**

- Milk. no withdrawal period

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

- Meat and offal. 47 day

Intramuscular use:

- **Cattle**
  - Milk. 7 day
  - Meat and offal. 48 day
- **Horse**
  - Milk. no withdrawal period

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

- Meat and offal. 47 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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Intervet Deutschland GmbH

Marketing authorisation date:
23/12/2005

Manufacturing sites for batch release:
Vet Pharma Friesoythe GmbH

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
6026382.00.00

Date of authorisation status change:
23/12/2005

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.