BORGAL LÖSUNG 200 mg/ml + 40 mg/ml Injektionslösung für Rinder, Schweine, Pferde, Meerschweinchen, Forellenbrut und Zierfische

Authorised

- Trimethoprim
- Sulfadoxine

Product identification

Medicine name:
BORGAL LÖSUNG 200 mg/ml + 40 mg/ml Injektionslösung für Rinder, Schweine, Pferde, Meerschweinchen, Forellenbrut und Zierfische

Active substance:
Trimethoprim
Sulfadoxine

Target species:
Guinea pig
Cattle
Horse
Pig
Ornamental fish
Trout (fry)

Route of administration:
Oral use
Intravenous use
Water-borne use
Subcutaneous use
Intramuscular use
Intratracheal use

Product details

Active substance and strength:
Trimethoprim
40.00 milligram(s) / 1.00 millilitre(s)

Sulfadoxine
200.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Cattle**
  - Meat and offal. 8 day
  - Milk. 4 day
- **Horse**
  - Milk. no withdrawal period

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

  - Meat and offal. 8 day

- **Pig**

- Meat and offal. 8 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 8 day
  - Milk. 4 day
- **Pig**
  - Meat and offal. 8 day

Intramuscular use:

- **Cattle**
  - Meat and offal. 8 day
  - Milk. 4 day
- **Horse**
  - Milk. no withdrawal period

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

- Meat and offal. 8 day

- **Pig**
  - Meat and offal. 8 day

Intratracheal use:

- **Cattle**
  - Meat and offal. 8 day
  - Milk. 4 day
-

Horse

- Milk. no withdrawal period

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

- Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01EW13

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

Authorisation status:
Valid

Authorised in:
Germany

Package description:
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:
Virbac

Marketing authorisation date:
11/08/2003

Manufacturing sites for batch release:

Intervet International GmbH
Virbac S.A.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
6489082.00.00

Date of authorisation status change:
11/08/2003

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.