Antastmon 500/100 mg/g Pulver zum Eingeben für Rinder (Kälber), Schweine (Läufer, Ferkel), Pferde (Fohlen), Schafe (Lämmer), Ziegen (Lämmer), Hunde

Authorised

- Sulfadiazine
- Trimethoprim

Product identification

Medicine name:
Antastmon 500/100 mg/g Pulver zum Eingeben für Rinder (Kälber), Schweine (Läufer, Ferkel), Pferde (Fohlen), Schafe (Lämmer), Ziegen (Lämmer), Hunde

Active substance:
Sulfadiazine
Trimethoprim

Target species:
Cattle (calf)
Dog
Goat (kid)
Sheep (lamb)
Horse (foal)
Pig (piglet)

Route of administration:
In-feed use
In drinking water use

Product details

Active substance and strength:
Sulfadiazine
500.00 milligram(s) / 1.00 gram(s)

Trimethoprim
100.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral powder

Withdrawal period by route of administration:
In-feed use:

- **Cattle (calf)**
  - Meat and offal. 10 day

- **Horse (foal)**
  - Meat and offal. 10 day

- **Pig (piglet)**
  - Meat and offal. 10 day

In drinking water use:

- **Horse (foal)**
  - Meat and offal. 10 day

- **Pig (piglet)**

- Meat and offal. 10 day

- **Cattle (calf)**
  - Meat and offal. 10 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01EW10

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
Available only in German
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:
Bela-Pharm GmbH & Co. KG

Marketing authorisation date:
24/09/2002

Manufacturing sites for batch release:
Bela-Pharm GmbH & Co. KG

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
6325386.00.00

Date of authorisation status change:
24/09/2002

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

Documents

Combined File of all Documents

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