Lotagen Konzentrat

Authorised

- Policresulen

Product identification

Medicine name:
Lotagen Konzentrat

Active substance:
Policresulen

Target species:
Cattle
Sheep
Pig
Dog
Horse

Route of administration:
Intracervical use
Cutaneous use
Intrauterine use
Intralesional use
Vaginal use

Product details

Active substance and strength:

Policresulen
360.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Concentrate for cutaneous solution

Withdrawal period by route of administration:

Intracervical use:

- **Cattle**
  - Meat and offal. 3 day
  - Milk. 1 day
- **Sheep**
  - Milk. 1 day
  - Meat and offal. 3 day
- **Pig**
  - Meat and offal. 3 day

Cutaneous use:

- **Cattle**
  - Meat and offal. 3 day
  - Milk. 1 day
- **Sheep**
  - Milk. 1 day
  - Meat and offal. 3 day
- **Horse**
  - Milk. 1 day

- Meat and offal. 3 day

- **Pig**
  - Meat and offal. 3 day

Intrauterine use:

- **Cattle**
  - Meat and offal. 3 day
  - Milk. 1 day

- **Sheep**
  - Milk. 1 day
  - Meat and offal. 3 day

- **Horse**
  - Milk. 1 day
  - Meat and offal. 3 day

- **Pig**
  - Meat and offal. 3 day

Intralesional use:

- **Cattle**
  - Meat and offal. 3 day
  - Milk. 1 day

- **Sheep**
  - Milk. 1 day
  - Meat and offal. 3 day

- **Horse**
  - Milk. 1 day
  - Meat and offal. 3 day
- **Pig**
  - Meat and offal. 3 day

Vaginal use:

- **Cattle**
  - Meat and offal. 3 day
  - Milk. 1 day
- **Sheep**
  - Milk. 1 day
  - Meat and offal. 3 day
- **Horse**
  - Milk. 1 day
  - Meat and offal. 3 day
- **Pig**
  - Meat and offal. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG51AD02

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

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:
19/04/2005

Manufacturing sites for batch release:
Trirx Segre

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
6174148.00.00

Date of authorisation status change:
19/04/2005

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

Documents

Summary of Product Characteristics

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Package Leaflet

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