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

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

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Sheep

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

Pig

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- Meat and offal. 3 day

Cutaneous use:

Cattle

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

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Dog

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Sheep

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

Horse

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

Pig

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- Meat and offal. 3 day

Intrauterine use:

Cattle

- Meat and offal. 3 day

- Milk. 1 day

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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

• Dog

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:

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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

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:

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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