

PROCAIN-PENICILLIN-G ad us.vet. 300 mg/ml Injektionssuspension für Rinder und Pferde

Authorised

- Benzylpenicillin procaine monohydrate

Product identification

Medicine name:

PROCAIN-PENICILLIN-G ad us.vet. 300 mg/ml Injektionssuspension für Rinder und Pferde

Active substance:

Benzylpenicillin procaine monohydrate

Target species:

Cattle
Horse

Route of administration:

Intramammary use
Intramuscular use

Product details

Active substance and strength:

Benzylpenicillin procaine monohydrate

300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramammary use:

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Cattle

- Milk. 6 day
- Meat and offal. 10 day für die Behandlungsdauer 3 Tage
- Meat and offal. 12 day 12 Tage für die Behandlungsdauer 4-7 Tage

Intramuscular use:

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Cattle

- Milk. 6 day
- Meat and offal. 10 day für die Behandlungsdauer 3 Tage
- Meat and offal. 12 day 12 Tage für die Behandlungsdauer 4-7 Tage

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Horse

- Milk. no withdrawal period

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

- Meat and offal. 10 day für die Behandlungsdauer 3 Tage
- Meat and offal. 12 day für die Behandlungsdauer 4-7 Tage

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CE09

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

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:

aniMedica GmbH

Marketing authorisation date:

1/08/2005

Manufacturing sites for batch release:

aniMedica GmbH

Industrial Veterinaria S.A.

aniMedica Herstellungs GmbH

Responsible authority:

Authorisation number:

6873219.00.00

Date of authorisation status change:

1/08/2005

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.