Alfacilline 15/15 LA suspensija injekcijām liellopiem, zirgiem, aitām, cūkām, suņiem un kaķiem

Authorised

- Benzylpenicillin procaine
- Benzathine benzylpenicillin

Product identification

Medicine name:
Alfacilline 15/15 LA suspensija injekcijām liellopiem, zirgiem, aitām, cūkām, suņiem un kaķiem

Active substance:
Benzylpenicillin procaine
Benzathine benzylpenicillin

Target species:
Dog
Cattle
Horse
Sheep
Pig
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Benzylpenicillin procaine
150.00 milligram(s) / 1.00 millilitre(s)

Benzathine benzylpenicillin
125.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 21 day
  - Milk. 6 day
- **Horse**
  - Meat and offal. 21 day
  - Milk. no withdrawal period

Nav reģistrēts lietošanai zirgiem, kuru pienu paredzēts izmantot cilvēku uzturā.

- **Sheep**
  - Meat and offal. 21 day
  - Milk. 6 day
- **Pig**
  - Meat and offal. 21 day

Subcutaneous use:

-

Sheep

- Meat and offal. 21 day
- Milk. 6 day

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Horse

- Meat and offal. 21 day
- Milk. no withdrawal period

Nav reģistrēts lietošanai zirgiem, kuru pienu paredzēts izmantot cilvēku uzturā.

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Cattle

- Meat and offal. 21 day
- Milk. 6 day

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Pig

- Meat and offal. 21 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CE30

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

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Authorisation status:
Valid

Authorised in:
Latvia

Package description:
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Alfasan International B.V.

Marketing authorisation date:
3/10/1997

Manufacturing sites for batch release:
Alfasan International B.V.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/97/0599

Date of authorisation status change:
5/10/1997

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.