

# Duplocillin LA, süstesuspensioon hobustele, veistele, sigadele, lammastele, koertele ja kassidele

Authorised

- Benzathine benzylpenicillin
- Benzylpenicillin procaine

## Product identification

### Medicine name:

Duplocillin LA, süstesuspensioon hobustele, veistele, sigadele, lammastele, koertele ja kassidele

### Active substance:

Benzathine benzylpenicillin  
Benzylpenicillin procaine

### Target species:

Cattle  
Horse (non-food-producing)  
Pig  
Sheep  
Dog  
Cat

### Route of administration:

Intramuscular use  
Subcutaneous use

## Product details

### Active substance and strength:

Benzathine benzylpenicillin

150000.00 international unit(s) / 1.00 millilitre(s)

Benzylpenicillin procaine

150000.00 international unit(s) / 1.00 millilitre(s)

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### Pharmaceutical form:

Suspension for injection

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### Withdrawal period by route of administration:

#### Intramuscular use:

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#### Cattle

- Meat and offal. 70 day
- Milk. 72 hour

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#### Horse (non-food-producing)

- Meat and offal. no withdrawal period

Mitte kasutada hobustel, kelle liha kavatsetakse tarvitada inимtoiduks.

- Milk. no withdrawal period

Mitte kasutada hobustel, kelle piima kavatsetakse tarvitada inимtoiduks.

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#### Pig

- Meat and offal. 70 day

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#### Sheep

- Meat and offal. 56 day
  - Milk. 72 hour
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01CE30

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Estonia

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**Package description:**

Available only in [Estonian](#)

Available only in [Estonian](#)

Available only in [Estonian](#)

Available only in [Estonian](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

5/02/2004

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**Manufacturing sites for batch release:**

Aprilia Animal Health S.r.l.

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**Responsible authority:**

State Agency Of Medicines

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**Authorisation number:**

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**Date of authorisation status change:**

5/02/2004

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

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