

Betamox LA, 150 mg/ml süstesuspensioon

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

Betamox LA, 150 mg/ml süstesuspensioon

Active substance:

Amoxicillin trihydrate

Target species:

Cattle

Pig

Sheep

Dog

Cat

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Amoxicillin trihydrate

172.20 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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Cattle

- Meat and offal. 39 day

- Milk. 108 hour
Piimale: 108 tundi (4,5 päeva).

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Pig

- Meat and offal. 42 day

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Sheep

- Meat and offal. 29 day

- Milk. no withdrawal period

Ei ole lubatud kasutamiseks lammastel, kelle piima tarvitatakse inimtoiduks.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Estonia

Available in:

Estonia

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

Legal basis of product authorisation:

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

Marketing authorisation holder:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

6/06/2002

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

State Agency Of Medicines

Authorisation number:

1053

Date of authorisation status change:

6/06/2002

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.