

Betamox Vet. injektionsvæske, suspension 150 mg/ml

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

Betamox Vet. injektionsvæske, suspension 150 mg/ml

Active substance:

Amoxicillin trihydrate

Target species:

Pig
Cattle
Cat
Sheep
Dog

Route of administration:

Subcutaneous use
Intramuscular 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:

Subcutaneous use:

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Pig

- Milk. 3 day
- Meat and offal. 30 day

•

Cattle

- Milk. 3 day
- Meat and offal. 30 day

•

Sheep

- Milk. 3 day
- Meat and offal. 30 day

Intramuscular use:

•

Pig

- Milk. 3 day
- Meat and offal. 30 day

•

Cattle

- Meat and offal. 30 day
- Milk. 3 day

•

Sheep

- Meat and offal. 30 day

- Milk. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Denmark

Package description:

Available only in [Danish](#)

Available only in [Danish](#)

Available only in [Danish](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Scanvet Animal Health A/S

Marketing authorisation date:

19/10/1988

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

Danish Medicines Agency

Authorisation number:

13229

Date of authorisation status change:

19/10/1988

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.