Noromox Prolongatum Vet. injektionsvæske, suspension 150 mg/ml

Authorised

- Amoxicillin trihydrate
- Amoxicillin trihydrate

Product identification

Medicine name:
Noromox Prolongatum Vet. injektionsvæske, suspension 150 mg/ml
Noromox Prolongatum Vet. 150 mg/ml injektionsvæske, suspension

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Active substance:
Amoxicillin trihydrate
Amoxicillin trihydrate

Target species:
Cat
Cattle
Sheep
Pig
Dog

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

Amoxicillin trihydrate
172.10 milligram(s) / 1.00 millilitre(s)

Amoxicillin trihydrate
172.10 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cat**
- **Cattle**
- **Sheep**
- **Pig**
- **Dog**

Intramuscular use:

- **Sheep**
- **Pig**
- **Dog**
- **Cat**
- **Cattle**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Denmark

Available in:
Denmark

Package description:
Available only in Danish
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:
30/10/1991

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Danish Medicines Agency

Authorisation number:
13590

Date of authorisation status change:
30/10/1991

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.
