

Multimox pulver og solvens til injektionsvæske, opløsning 212 mg/ml

Authorised

- Amoxicillin sodium

Product identification

Medicine name:

Multimox pulver og solvens til injektionsvæske, opløsning 212 mg/ml

Active substance:

Amoxicillin sodium

Target species:

Pig

Cattle

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Amoxicillin sodium

212.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Powder and solvent for solution for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Pig

- Meat and offal. 9 day

-

Cattle

- Milk. 3 day

- Meat and offal. 9 day

Intravenous use:

-

Cattle

- Meat and offal. 5 day

- Milk. 24 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

Available only in Danish

Available only in Danish

Additional information

Entitlement type:

Marketing Authorisation

Marketing authorisation holder:

Scanvet Animal Health A/S

Marketing authorisation date:

2/11/2017

Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

Responsible authority:

Danish Medicines Agency

Authorisation number:

58257

Date of authorisation status change:

2/11/2017

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.