

Maxoject LA 200 mg/ml Solution for Injection

Authorised

- Oxytetracycline

Product identification

Medicine name:

Maxoject LA 200 mg/ml Solution for Injection

Active substance:

Oxytetracycline

Target species:

Cattle

Sheep

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Oxytetracycline

200.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

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Cattle

- Meat and offal. 35 day
- Milk. 8 day

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Sheep

- Meat and offal. 20 day
- Milk. 8 day

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Pig

- Meat and offal. 20 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Amber type II glass vials of 100 ml with nitryl bungs and aluminium seals

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:

Chem-Pharm (Ballyvaughan) Limited

Marketing authorisation date:

1/10/1998

Manufacturing sites for batch release:

Norbrook Laboratories Limited

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10823/007/001

Date of authorisation status change:

1/10/1998

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics