MACROSYN 100 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS AND SHEEP

Authorised

- Tulathromycin

Product identification

Medicine name:
MACROSYN 100 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS AND SHEEP
Macrosyn 100 mg/ml Oplossing voor injectie
Macrosyn 100 mg/ml Solution injectable
Macrosyn 100 mg/ml Injektionslösung

Active substance:
Tulathromycin

Target species:
Sheep
Pig
Cattle

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Tulathromycin
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Sheep**

  - Meat and offal. 16 day

  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption. Do not use in pregnant animals, which are intended to produce milk for human consumption, within 2 months of expected parturition.

- **Pig**

  - Meat and offal. 13 day

Subcutaneous use:

- **Cattle**

  - Meat and offal. 22 day

  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption. Do not use in pregnant animals, which are intended to produce milk for human consumption, within 2 months of expected parturition.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01FA94

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bimeda Animal Health Limited

Marketing authorisation date:
17/09/2020

Manufacturing sites for batch release:
Bimeda Animal Health Limited

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V571146

Date of authorisation status change:
17/09/2020

Reference member state:
France

Procedure number:
FR/V/0418/001

Concerned member states:
Austria Belgium Denmark Estonia Germany Ireland Italy Latvia Lithuania Netherlands Poland Spain

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

Documents

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

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