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# 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 SOLUCION INYECTABLE PARA BOVINO, PORCINO Y OVINO

#### **Active substance:**

Tulathromycin

# **Target species:**

Sheep

Pig

Cattle

#### Route of administration:

Intramuscular use

Subcutaneous use

# **Product details**

# **Active substance and strength:**

#### **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:**

#### **Authorised in:**

Spain

## 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:

9/09/2020

# Manufacturing sites for batch release:

Bimeda Animal Health Limited

## **Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

#### **Authorisation number:**

3935 ESP

## Date of authorisation status change:

10/09/2020

#### 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

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

## **Documents**

Labelling

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

Summary of Product Characteristics

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

Package Leaflet

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