

PULMOXXIN 100 MG/ML SOLUTION INJECTABLE POUR BOVINS PORCINS ET OVINS

Authorised

- Tulathromycin

Product identification

Medicine name:

PULMOXXIN 100 MG/ML SOLUTION INJECTABLE POUR BOVINS PORCINS ET OVINS

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

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine. Ne pas utiliser chez les animaux gravides producteurs de lait destiné à la consommation humaine au cours des 2 mois précédant la mise bas.

-

Pig

- Meat and offal. 13 day

Subcutaneous use:

-

Cattle

- Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine. Ne pas utiliser chez les animaux gravides producteurs de lait destiné à la consommation humaine au cours des 2 mois précédant la mise bas.

- Meat and offal. 22 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA94

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

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:

Laboratoires Biove

Marketing authorisation date:

28/01/2021

Manufacturing sites for batch release:

Laboratoires Biove

V.M.D.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/6204909 8/2021

Date of authorisation status change:

28/01/2021

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.

Package Leaflet and Labelling

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

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