

IVERTIN 10 mg/ml Solution for Injection for Cattle and Pigs

Authorised

- Ivermectin

Product identification

Medicine name:

IVERTIN 10 mg/ml Solution for Injection for Cattle and Pigs

Active substance:

Ivermectin

Target species:

Cattle

Pig

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Ivermectin

10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

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

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Cattle

- Milk. no withdrawal period

Do not use in lactating dairy cows producing milk for human consumption. Do not use in non-lactating dairy cows including pregnant heifers within 60 days of calving.

- Meat and offal. 49 day

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Pig

- Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

(ID12) 6000 millilitre(s): unspecified outer container with 12 Vial (polypropylene) each with 500 millilitre(s)

(ID11) 1200 millilitre(s): unspecified outer container with 12 Vial (polypropylene) each with 100 millilitre(s)

(ID10) 600 millilitre(s): unspecified outer container with 12 Vial (polypropylene) each with 50 millilitre(s)

(ID9) 5000 millilitre(s): unspecified outer container with 10 Vial (polypropylene) each with 500 millilitre(s)

(ID8) 1000 millilitre(s): unspecified outer container with 10 Vial (polypropylene) each with 100 millilitre(s)

(ID7) 500 millilitre(s): unspecified outer container with 10 Vial (polypropylene) each with 50 millilitre(s)

(ID6) 3000 millilitre(s): unspecified outer container with 6 Vial (polypropylene) each with 500 millilitre(s)

(ID5) 600 millilitre(s): unspecified outer container with 6 Vial (polypropylene) each with 100 millilitre(s)

(ID4) 300 millilitre(s): unspecified outer container with 6 Vial (polypropylene) each with 50 millilitre(s)

(ID3) 500 millilitre(s): unspecified outer container with 1 Vial (polypropylene) with 500 millilitre(s)

(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (polypropylene) with 100 millilitre(s)

(ID1) 50 millilitre(s): unspecified outer container with 1 Vial (polypropylene) with 50 millilitre(s)

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:

Laboratorios Calier S.A.

Marketing authorisation date:

29/09/2005

Manufacturing sites for batch release:

Laboratorios Calier S.A.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/3943503 9/2005

Date of authorisation status change:

22/12/2021

Reference member state:

Germany

Procedure number:

DE/V/0165/001

Concerned member states:

Austria Belgium France Italy United Kingdom (Northern Ireland)

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.

Combined File of all Documents