Zuprevo 180 mg/ml - Solution for injection

Authorised

- Tildipirosin

Product identification

Medicine name:
Zuprevo 180 mg/ml - Solution for injection

Active substance:
Tildipirosin

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Tildipirosin
180.00 milligram(s) / 1.00 Vial

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 47 day

47 days (Not authorised for use in lactating 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:

QJ01FA96

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Croatia , Czechia , Denmark , France , Germany , Ireland , Italy , Luxembourg , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , United Kingdom (Northern Ireland)

Package description:

Packaging:Vial (glass), Package_size:1 vial, Content:250 ml

Packaging:Vial (glass), Package_size:1 vial, Content:100 ml

Packaging:Vial (glass), Package_size:1 vial, Content:50 ml

Packaging:Vial (glass), Package_size:1 vial, Content:20 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
6/05/2011

Manufacturing sites for batch release:
Intervet International GmbH

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
6/05/2011

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

Documents

Combined File of all Documents

English (PDF)

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