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ECOMECTIN 10 mg/ml injekčný roztok

Authorised

- Ivermectin

Product identification

Medicine name:
ECOMECTIN 10 mg/ml injekčný roztok

Active substance:
Ivermectin

Target species:
Cattle
Sheep
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:

- **Cattle**
  - Meat and offal. 49 day

Not authorised for use in animals producing milk for human consumption.

- **Sheep**
  - Meat and offal. 42 day

Not authorised for use in animals producing milk for human consumption.

- **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:

Slovakia

Package description:

Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Eco Animal Health Europe Limited

Marketing authorisation date:
2/05/2002

Manufacturing sites for batch release:
Divasa Farmavic S.A.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/027/02-S

Date of authorisation status change:
2/05/2002

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

Documents

Combined File of all Documents

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