Bremamectin, 10 mg/ml süstelahus veistele, lammastele, kitsedele ja sigadele

Authorised

- Ivermectin

Product identification

Medicine name:
Bremamectin, 10 mg/ml süstelahus veistele, lammastele, kitsedele ja sigadele

Active substance:
Ivermectin

Target species:
Cattle
Pig
Sheep
Goat

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
  - Milk. no withdrawal period

Ei ole lubatud kasutamiseks loomadel, kelle piima tarvitatakse inimtoiduks. Mitte kasutada tiinetel loomadel, kelle piima kavatsetakse tarvitada inimtoiduks, 33 päeva enne oodatavat poegimist.

- **Pig**
  - Meat and offal. 28 day

- **Sheep**
  - Meat and offal. 28 day
  - Milk. no withdrawal period

Ei ole lubatud kasutamiseks loomadel, kelle piima tarvitatakse inimtoiduks. Mitte kasutada tiinetel loomadel, kelle piima kavatsetakse tarvitada inimtoiduks, 33 päeva enne oodatavat poegimist.

- **Goat**
  - Meat and offal. 28 day
  - Milk. no withdrawal period

Ei ole lubatud kasutamiseks loomadel, kelle piima tarvitatakse inimtoiduks. Mitte kasutada tiinetel loomadel, kelle piima kavatsetakse tarvitada inimtoiduks, 33 päeva enne oodatavat poegimist.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Estonia

Package description:
Available only in Estonian
Available only in Estonian
Available only in Estonian

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:
Bremer Pharma GmbH

Marketing authorisation date:
17/08/2001

Manufacturing sites for batch release:
Laboratorios Karizoo S.A.

Responsible authority:
State Agency Of Medicines

Authorisation number:
1002

Date of authorisation status change:
17/08/2001

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

Documents

Summary of Product Characteristics

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