FERROFERON 200 mg/ml Solution for injection for pigs

Not authorised

- Gleptoferron

Product identification

Medicine name:
FERROFERON 200 mg/ml Solution for injection for pigs
Ursoglepto Vet. 200 mg/ml injektionsvæske, opløsning

Active substance:
Gleptoferron

Target species:
Pig (sucking piglet)

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Gleptoferron
532.60 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Pig (sucking piglet)**

  - Meat and offal. 0 day

Intramuscular use:

- **Pig (sucking piglet)**

  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QB03AC

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

---

Authorisation status:
Surrendered

Authorised in:
Denmark

Package description:
(ID4) 1000 millilitre(s): unspecified outer container with 10 Vial (Low Density PolyEthylene) each with 100 millilitre(s)
(ID6) 2000 millilitre(s): unspecified outer container with 10 Vial (Low Density PolyEthylene) each with 200 millilitre(s)
(ID5) 200 millilitre(s): unspecified outer container with 1 Vial (Low Density PolyEthylene) with 200 millilitre(s)
(ID3) 100 millilitre(s): unspecified outer container with 1 Vial (Low Density PolyEthylene) with 100 millilitre(s)
(ID2) 1000 millilitre(s): unspecified outer container with 10 Vial (Glass) each with 100 millilitre(s)
(ID1) 100 millilitre(s): unspecified outer container with 1 Vial (Glass) with 100 millilitre(s)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:
Iron4u ApS

Marketing authorisation date:
19/08/2013

Manufacturing sites for batch release:
Ceva Sante Animale
Serumwerk Bernburg AG

Responsible authority:
Danish Medicines Agency

Authorisation number:
52519

Date of authorisation status change:
13/05/2024

Reference member state:
Germany

Procedure number:
DE/V/0157/001

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 24/03/2023
Download
