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
Ferroferon 200 mg/ml

Active substance:
Gleptoferron

Target species:
Pig (suckling 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 (suckling piglet)**
  - Meat and offal. 0 day

Intramuscular use:

- **Pig (suckling piglet)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QB03AC

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Germany

Package description:

(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)
(ID4) 1000 millilitre(s): unspecified outer container with 10 Vial (Low Density PolyEthylene) each with 100 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:
4/08/2011

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

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401405.00.00

Date of authorisation status change:
15/05/2024

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

Package Leaflet

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