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

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

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Authorisation status:
Surrendered

Authorised in:
Belgium

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

Marketing authorisation date:
21/01/2014

Manufacturing sites for batch release:
CEVA SANTE ANIMALE - LIBOURNE
Serumwerk Bernburg AG

Responsible authority:
Federal Agency For Medicines And Health Products

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

Date of authorisation status change:
20/06/2022

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

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.