

# 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

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**Active substance:**

Gleptoferron

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**Target species:**

Pig (suckling piglet)

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**Route of administration:**

Subcutaneous use  
Intramuscular use

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## Product details

**Active substance and strength:**

Gleptoferron  
532.60 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Subcutaneous use:**

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**Pig (suckling piglet)**

- Meat and offal. 0 day

**Intramuscular use:**

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**Pig (suckling piglet)**

- Meat and offal. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QB03AC

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**Legal status of supply:**

Veterinary medicinal product not subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

Denmark

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**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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Informed consent application (Article 13c of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Iron4u ApS

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**Marketing authorisation date:**

19/08/2013

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**Manufacturing sites for batch release:**

Ceva Sante Animale

Serumwerk Bernburg AG

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**Responsible authority:**

Danish Medicines Agency

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**Authorisation number:**

52519

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**Date of authorisation status change:**

13/05/2024

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0157/001

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

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