# URSOFERRAN 200 mg/ml Solution for injection for pigs

Authorised

- Gleptoferron

## Product identification

**Medicine name:**
URSOFERRAN 200 mg/ml Solution for injection for pigs
Ursoferran 200 mg/ml oplossing voor injectie voor varkens

**Active substance:**
Gleptoferron

**Target species:**
Pig (sucking piglet)

**Route of administration:**
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:**

**Intramuscular use:**

- **Pig (sucking 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 subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Netherlands

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**Available in:**

Netherlands

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**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 10 Vial (Glass) with 100 millilitre(s) (1000 millilitre(s))
(ID2): 1 unspecified outer container with 1 Vial (Low Density PolyEthylene) with 100 millilitre(s) (100 millilitre(s))
(ID1): 1 unspecified outer container with 1 Vial (Glass) with 100 millilitre(s) (100 millilitre(s))

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**
Well-established use application (Article 13a of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Serumwerk Bernburg AG

**Marketing authorisation date:**
15/04/2013

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

**Responsible authority:**
Medicines Evaluation Board

**Authorisation number:**
REG NL 113079

**Date of authorisation status change:**
8/02/2022

**Reference member state:**
Germany

**Procedure number:**
DE/V/0149/001

**Concerned member states:**
Belgium Czechia Denmark Finland France Hungary Italy Luxembourg Netherlands Poland Portugal Romania Slovakia

United Kingdom (Northern Ireland)

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

## Documents

Summary of Product Characteristics

Combined File of all Documents

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