

URSOFERRAN 200 mg/ml Solution for injection for pigs

Not
authorised

- Gleptoferron

Product identification

Medicine name:

URSOFERRAN 200 mg/ml Solution for injection for pigs

URSOFERRAN 200 mg/ml injekčný roztok pre ošípané

Active substance:

Gleptoferron

Target species:

Pig (suckling 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 (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:

Slovakia

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))

Additional information

Entitlement type:

Marketing Authorisation

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:

20/05/2013

Manufacturing sites for batch release:

Serumwerk Bernburg AG

Ceva Sante Animale

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/031/MR/13-S

Date of authorisation status change:

4/10/2022

Reference member state:

Germany

Procedure number:

DE/V/0149/001

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

Documents

Combined File of all Documents

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