URSOFERRAN 200 mg/ml Solution for injection for pigs

Authorised

- Gleptoferron

Product identification

Medicine name:
URSOFERRAN 200 mg/ml Solution for injection for pigs

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

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Authorisation status:

Valid

Authorised in:

Germany

Available 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) 1000 millilitre(s): unspecified outer container with 10 Vial (glass) each with 100 millilitre(s)
(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (low-density polyethylene) 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:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Serumwerk Bernburg AG

Marketing authorisation date:
22/08/2005

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

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
3100349.00.00

Date of authorisation status change:
13/12/2017

Reference member state:
Germany

Procedure number:
DE/V/0149/001

Concerned member states:
Belgium Czechia Denmark Finland France Hungary Italy Netherlands Romania Slovakia United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
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