URSOFERRAN 200 mg/ml Solution for injection for pigs

Authorised

- Gleptoferron

Product identification

Medicine name:
URSOFERRAN 200 mg/ml Solution for injection for pigs
Viloferron 200 mg/ml injektionsvæske, opløsning

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QB03AC

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Denmark

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:
19/08/2013

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

Responsible authority:
Danish Medicines Agency

Authorisation number:
51985

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

English (PDF)

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