Hypophysin LA 35 µg/ml solution for injection for cattle and pigs

Authorised

- Carbetocin

Product identification

Medicine name:
Hypophysin LA 35 µg/ml solution for injection for cattle and pigs
Depotocin 35 µg/ml Injektionslösung für Rinder und Schweine

Active substance:
Carbetocin

Target species:
Cattle
Pig

Route of administration:
Intravenous use
Intramuscular use

Product details

Active substance and strength:
Carbetocin
35.00 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Intramuscular use:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH01BB03

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:

(ID5) 100 millilitre(s): unspecified outer container with 1 Vial (Glass) with 100 millilitre(s)
(ID4) 600 millilitre(s): unspecified outer container with 12 Vial (Glass) each with 50 millilitre(s)
(ID3) 50 millilitre(s): unspecified outer container with 1 Vial (Glass) with 50 millilitre(s)
(ID2) 20 millilitre(s): unspecified outer container with 1 Vial (Glass) with 20 millilitre(s)
(ID1) 10 millilitre(s): unspecified outer container with 1 Vial (Glass) with 10 millilitre(s)
(ID6) 1200 millilitre(s): unspecified outer container with 12 Vial (Glass) each with 100 millilitre(s)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Veyx Pharma GmbH

Marketing authorisation date:
26/05/2014

Manufacturing sites for batch release:
Veyx Pharma GmbH

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401958.00.00

Date of authorisation status change:
29/07/2019

Reference member state:
Germany

Procedure number:
DE/V/0156/001

Concerned member states:
Austria Belgium Bulgaria Czechia Estonia France Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

English (RTF)
Published on: 3/02/2025
Download

Summary of Product Characteristics

English (RTF)
Published on: 3/02/2025
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