Hypophysin LA 35 microgram/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
Hypophysin LA 35 microgram/ml Solution for Injection for Cattle and Pigs

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

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

Package description:
(ID5) 100 millilitre(s): unspecified outer container with 1 Vial (glass) with 100 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)

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:
5/08/2014

Manufacturing sites for batch release:
Veyx Pharma GmbH
Veyx-Pharma B.V.

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 27569/4003

Date of authorisation status change:
1/10/2020

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