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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

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

Intramuscular use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 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:

Lithuania

Available in:

Lithuania

Package description:
(ID5) 100 millilitre(s): Box (Cardboard) with 1 Vial (Glass type I) with 100 millilitre(s), closed with Cap (Aluminium) and Stopper (Bromobutyl Rubber)
(ID3) 50 millilitre(s): Box (Cardboard) with 1 Vial (Glass type I) with 50 millilitre(s), closed with Cap (Aluminium) and Stopper (Bromobutyl Rubber)
(ID2) 20 millilitre(s): Box (Cardboard) with 1 Vial (Glass type I) with 20 millilitre(s), closed with Cap (Aluminium) and Stopper (Bromobutyl Rubber)
(ID1) 10 millilitre(s): Box (Cardboard) with 1 Vial (Glass type I) with 10 millilitre(s), closed with Cap (Aluminium) and Stopper (Bromobutyl Rubber)

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:
25/06/2014

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

Responsible authority:
State Food And Veterinary Service

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
28/06/2026

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

Combined File of all Documents

RV2231.pdf

2401958-paren-20140521.pdf