

# Hypophysin LA 70 µg/ml solution for injection for cattle and pigs

Authorised

- Carbetocin

## Product identification

**Medicine name:**

Hypophysin LA 70 µg/ml solution for injection for cattle and pigs  
Depotocin 70 µg/ml Injektionslösung für Rinder und Schweine

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**Active substance:**

Carbetocin

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**Target species:**

Cattle  
Pig

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**Route of administration:**

Intravenous use  
Intramuscular use

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## Product details

**Active substance and strength:**

Carbetocin  
70.00 microgram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intravenous use:****• Cattle**

- Meat and offal. 0 day
- Milk. 0 hour

**• Pig**

- Meat and offal. 0 day

**Intramuscular use:****• Cattle**

- Meat and offal. 0 day
- Milk. 0 hour

**• Pig**

- Meat and offal. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QH01BB03

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Germany

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**Available in:**

Germany

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**Package description:**

(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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Veyx Pharma GmbH

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**Marketing authorisation date:**

26/05/2014

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**Manufacturing sites for batch release:**

Veyx Pharma GmbH

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

401959.00.00

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**Date of authorisation status change:**

29/07/2019

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0156/002

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**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)

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

Package Leaflet

2401959-paren-20140521.pdf

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**Source URL:** <https://medicines.health.europa.eu/veterinary/600000061033>