

# DEPOTOCIN, 0.07mg/ml, Injekční roztok

Not  
authorised

- Carbetocin

## Product identification

**Medicine name:**

DEPOTOCIN, 0.07mg/ml, Injekční roztok

**Active substance:**

Carbetocin

**Target species:**

Dog (bitch)

Cattle (cow)

Goat

Sheep

Pig (sow)

**Route of administration:**

Intravenous use

Subcutaneous use

Intramuscular use

## Product details

**Active substance and strength:**

Carbetocin

0.07 milligram(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:**

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**Cattle (cow)**

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

- 

**Goat**

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

- 

**Sheep**

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

- 

**Pig (sow)**

- Meat and offal. 0 day

**Subcutaneous use:**

- 

**Cattle (cow)**

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

- 

**Goat**

- Meat and offal. 0 day

- Milk. 0 day

- 

**Sheep**

- Meat and offal. 0 day

- Milk. 0 day

- 

**Pig (sow)**

- Meat and offal. 0 day

**Intramuscular use:**

- 

**Cattle (cow)**

- Meat and offal. 0 day

- Milk. 0 day

- 

**Goat**

- Meat and offal. 0 day

- Milk. 0 day

- 

**Sheep**

- Meat and offal. 0 day

- Milk. 0 day

- 

**Pig (sow)**

- 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:**

Surrendered

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**Authorised in:**

Czechia

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**Package description:**

Available only in Czech

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Vetoquinol S.A.

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**Marketing authorisation date:**

24/06/1992

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**Manufacturing sites for batch release:**

NORDIC Pharma s.r.o.

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/415/92-C

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**Date of authorisation status change:**

25/09/2024

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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

This document does not exist in this language (English). You can find it in another language below.

### Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

### Labelling

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