

## Lactovac C

Not  
authorised

- Escherichia coli, serotype O9:K35 (fimbrial adhesin F5 and F41), strain S1091/83, Inactivated
- Bovine rotavirus A, strain 1005/78, Inactivated
- Bovine rotavirus A, strain Holland, Inactivated
- Bovine coronavirus, strain 800, Inactivated

## Product identification

**Medicine name:**

Lactovac C

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**Active substance:**

Escherichia coli, serotype O9:K35 (fimbrial adhesin F5 and F41), strain S1091/83, Inactivated

Bovine rotavirus A, strain 1005/78, Inactivated

Bovine rotavirus A, strain Holland, Inactivated

Bovine coronavirus, strain 800, Inactivated

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**Target species:**

Cattle

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**Route of administration:**

Subcutaneous use

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## Product details

### **Active substance and strength:**

Escherichia coli, serotype O9:K35 (fimbrial adhesin F5 and F41), strain S1091/83, Inactivated

1.00 relative potency / 5.00 millilitre(s)

Bovine rotavirus A, strain 1005/78, Inactivated

1.00 relative potency / 5.00 millilitre(s)

Bovine rotavirus A, strain Holland, Inactivated

1.00 relative potency / 5.00 millilitre(s)

Bovine coronavirus, strain 800, Inactivated

1.00 relative potency / 5.00 millilitre(s)

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### **Pharmaceutical form:**

Suspension for injection

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### **Withdrawal period by route of administration:**

#### **Subcutaneous use:**

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#### **Cattle**

- Meat and offal. 0 day

- Milk. 0 day

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

QI02AL01

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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:**

Germany

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### **Package description:**

Available only in [German](#)

Available only in [German](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Zoetis Deutschland GmbH

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**Marketing authorisation date:**

6/04/1999

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**Manufacturing sites for batch release:**

Zoetis Belgium

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**Responsible authority:**

Paul-Ehrlich-Institut

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**Authorisation number:**

PEI.V.00747.01.1

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**Date of authorisation status change:**

9/08/2022

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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

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