

# Rispoval IBR-Marker InactivatumSuspension for injection for cattle

Not  
authorised

- Bovine herpesvirus 1, strain Difivac gE gene-deleted, Inactivated

## Product identification

### **Medicine name:**

Rispoval IBR-Marker InactivatumSuspension for injection for cattle

Rispoval IBR-Marker Inactivatum, суспензия за инжектиране за говеда

### **Active substance:**

Bovine herpesvirus 1, strain Difivac gE gene-deleted, Inactivated

### **Target species:**

Cattle

Cattle (for meat production)

Cattle (calf)

Cattle (heifer)

### **Route of administration:**

Subcutaneous use

## Product details

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

Bovine herpesvirus 1, strain Difivac gE gene-deleted, Inactivated  
0.01 titre / 2.00 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**

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

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#### **Cattle (for meat production)**

- Meat and offal. 0 day

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#### **Cattle (calf)**

- Meat and offal. 0 day

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#### **Cattle (heifer)**

- Meat and offal. 0 day
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### **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI02AA03

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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:**

Bulgaria

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**Package description:**

(ID3) 100 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 100 millilitre(s)  
(ID2) 20 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 20 millilitre(s)

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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 Belgium

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**Marketing authorisation date:**

9/07/2008

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**Manufacturing sites for batch release:**

Zoetis Belgium

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**Responsible authority:**

Bulgarian Food Safety Authority

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**Authorisation number:**

0022-1295-01.02.2010

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**Date of authorisation status change:**

16/04/2024

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0021/001

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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 and Labelling

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

### Combined File of all Documents