

# BIOSUIS Salm emulsion for injection for pigs

Authorised

- Salmonella enterica, subsp. enterica, serovar Typhimurium, Inactivated
- Salmonella enterica, subsp. enterica, serovar Derby, Inactivated
- Salmonella enterica, subsp. enterica, serovar Infantis, Inactivated

## Product identification

**Medicine name:**

BIOSUIS Salm emulsion for injection for pigs

---

**Active substance:**

Salmonella enterica, subsp. enterica, serovar Typhimurium, Inactivated

Salmonella enterica, subsp. enterica, serovar Derby, Inactivated

Salmonella enterica, subsp. enterica, serovar Infantis, Inactivated

---

**Target species:**

Pig (pregnant sow)

Pig (pregnant gilt)

---

**Route of administration:**

Intramuscular use

---

## Product details

### Active substance and strength:

Salmonella enterica, subsp. enterica, serovar Typhimurium, Inactivated

1.00 relative potency / 1.00 Dose

Salmonella enterica, subsp. enterica, serovar Derby, Inactivated

1.00 relative potency / 1.00 Dose

Salmonella enterica, subsp. enterica, serovar Infantis, Inactivated

1.00 relative potency / 1.00 Dose

---

### Pharmaceutical form:

Emulsion for injection

---

### Withdrawal period by route of administration:

#### Intramuscular use:

- 

#### **Pig (pregnant sow)**

- Meat and offal. 0 day

- 

#### **Pig (pregnant gilt)**

- Meat and offal. 0 day

---

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

QI09AB14

---

### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

### Authorisation status:

Valid

---

### Authorised in:

Netherlands

---

### Package description:

Glass Vial 1 x 10.0 Dose

Plastic Vial 1 x 50.0 Dose  
Plastic Vial 1 x 100.0 Dose  
Glass Vial 10 x 10.0 Dose

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

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

---

**Marketing authorisation holder:**

Bioveta a.s.

---

**Marketing authorisation date:**

8/11/2019

---

**Manufacturing sites for batch release:**

Bioveta a.s.

---

**Responsible authority:**

Medicines Evaluation Board

---

**Authorisation number:**

REG NL 123500

---

**Date of authorisation status change:**

28/01/2022

---

**Reference member state:**

Czechia

---

**Procedure number:**

CZ/V/0151/001

---

**Concerned member states:**

Austria Belgium Estonia France Germany Greece Hungary Ireland Italy  
Netherlands Poland Slovakia United Kingdom (Northern Ireland)

---

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

## Documents

Combined File of all Documents

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

Summary of Product Characteristics

Package Leaflet

Labelling

eu-puar-czv0151001-mr-biosuis\_salm-en.pdf