

# Salmoporc, lyophilisate and solvent for suspension for injection for pigs

Authorised

- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 421/125 (histidine-adenine auxotrophic), Live

## Product identification

**Medicine name:**

Salmoporc, lyophilisate and solvent for suspension for injection for pigs

---

**Active substance:**

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 421/125 (histidine-adenine auxotrophic), Live

---

**Target species:**

Pig

---

**Route of administration:**

Oral use

Subcutaneous use

---

## Product details

**Active substance and strength:**

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 421/125 (histidine-adenine auxotrophic), Live

8.00 log<sub>10</sub> colony forming unit(s) / 1.00 millilitre(s)

---

**Pharmaceutical form:**

Lyophilisate for suspension for injection

---

**Withdrawal period by route of administration:**

**Oral use:**

- 

**Pig**

- Meat and offal. 6 week  
six weeks after the second vaccination

**Subcutaneous use:**

- 

**Pig**

- Meat and offal. 6 week  
six weeks after the second vaccination

---

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

QI09AE02

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Slovakia

---

**Package description:**

Cardboard box containing 1 vial with 20 doses (1 dose = 1 ml) lyophilised vaccine  
1 vial with 20 ml solvent

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Similar biological application (Article 13(4) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Ceva Animal Health Slovakia s.r.o.

---

**Marketing authorisation date:**

7/06/2019

---

**Manufacturing sites for batch release:**

Ceva-Phylaxia Zrt.

IDT Biologika GmbH

---

**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

---

**Authorisation number:**

97/004/DC/19-S

---

**Date of authorisation status change:**

7/06/2019

---

**Reference member state:**

Netherlands

---

**Procedure number:**

NL/V/0247/001

---

**Concerned member states:**

Austria Belgium Czechia Denmark Hungary Ireland Italy Portugal Romania  
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.