

# 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

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**Active substance:**

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

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**Target species:**

Pig

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**Route of administration:**

Oral use

Subcutaneous use

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## 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)

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**Pharmaceutical form:**

Lyophilisate for suspension for injection

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**Withdrawal period by route of administration:**

**Oral use:**

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**Pig**

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

**Subcutaneous use:**

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**Pig**

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

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

QI09AE02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Romania

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**Package description:**

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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Ceva Sante Animale Romania S.R.L.

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**Marketing authorisation date:**

31/03/2019

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**Manufacturing sites for batch release:**

Ceva-Phylaxia Zrt.

IDT Biologika GmbH

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**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

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**Authorisation number:**

240045

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**Date of authorisation status change:**

26/05/2026

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0247/001

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**Concerned member states:**

Austria Belgium Czechia Denmark Hungary Ireland Italy Portugal Romania  
Slovakia United Kingdom (Northern Ireland)

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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

Combined File of all Documents

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