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# 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 log10 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:**
Ireland

**Available in:**
Ireland

**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 Sante Animale

**Marketing authorisation date:**
18/04/2019

**Manufacturing sites for batch release:**
Ceva-Phylaxia Zrt.
IDT Biologika GmbH

**Responsible authority:**
Health Products Regulatory Authority

**Authorisation number:**
VPA10815/064/001

**Date of authorisation status change:**
18/04/2019

**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)

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