

# SALMOPORC SCS lyophilisate and solvent for suspension for pigs

Not authorised

- Salmonella enterica, subsp. enterica, serovar Choleraesuis, strain 431/261, Live

## Product identification

**Medicine name:**

SALMOPORC SCS lyophilisate and solvent for suspension for pigs

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**Active substance:**

Salmonella enterica, subsp. enterica, serovar Choleraesuis, strain 431/261, Live

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**Target species:**

Pig

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**Route of administration:**

Subcutaneous use

Oral use

Intramuscular use

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## Product details

**Active substance and strength:**

Salmonella enterica, subsp. enterica, serovar Choleraesuis, strain 431/261, Live

8273.00 [CFU] /dose / 1.00 [CFU] /dose

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**Pharmaceutical form:**

Lyophilisate and solvent for emulsion for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Pig**

- Meat and offal. 3 week Meat and offal: 3 weeks

**Oral use:**

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**Pig**

- Meat and offal. 3 week Meat and offal: 3 weeks

**Intramuscular use:**

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**Pig**

- Meat and offal. 3 week Meat and offal: 3 weeks

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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:**

Surrendered

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**Authorised in:**

Germany

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**Package description:**

Lyophilised vaccine: 10 ml hydrolytic class 1 glass bottles, each containing 50 doses of vaccine and closed with bromobutyl rubber stopper and secured caps. Solvent: 50 ml hydrolytic class 2 glass bottles, each containing 50 ml of solvent, closed with

bromobutyl rubber stopper and secured caps. Cardboard carton containing 50 doses (lyophilisate and solvent).

Lyophilised vaccine: 10 ml hydrolytic class 1 glass bottles, each containing 10 doses of vaccine and closed with bromobutyl rubber stopper and secured caps. Solvent: 10 ml hydrolytic class 2 glass bottles, each containing 10 ml of solvent, closed with bromobutyl rubber stopper and secured caps. Cardboard carton containing 10 doses (lyophilisate and solvent).

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Ceva Tiergesundheit GmbH

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**Marketing authorisation date:**

23/01/2017

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**Manufacturing sites for batch release:**

IDT Biologika GmbH

Solupharm Pharmazeutische Erzeugnisse GmbH

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**Responsible authority:**

Paul-Ehrlich-Institut

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**Authorisation number:**

PEI.V.11873.01.1

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**Date of authorisation status change:**

20/04/2023

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**Reference member state:**

Poland

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**Procedure number:**

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.