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
Suisaloral

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

Target species:
Pig

Route of administration:
Subcutaneous use
Oral use
Intramuscular use

Product details

Active substance and strength:
Salmonella enterica, subsp. enterica, serovar Choleraesuis, strain 431/261, Live
8273.00 colony forming unit(s)/dose / 1.00 colony forming unit(s)/dose

Pharmaceutical form:
Lyophilisate and solvent for emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Pig**

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

Oral use:

- **Pig**

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

Intramuscular use:

- **Pig**

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AE02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Surrendered

Authorised in:
Germany

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).

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:
Ceva Tiergesundheit GmbH

Marketing authorisation date:
23/01/2017

Manufacturing sites for batch release:
IDT Biologika GmbH
Solupharm Pharmazeutische Erzeugnisse GmbH

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.11873.01.1

Date of authorisation status change:
15/05/2023

Reference member state:
Poland

Procedure number:
PL/V/0106/001/MR

To consult adverse reactions on veterinary medicinal products please go to 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.
