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

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 [CFU] /dose / 1.00 [CFU] /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:
This information is not available for this product.

Authorisation status:
Revoked

Authorised in:
Poland

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 Animal Health Polska Sp. z o.o.

Marketing authorisation date:
20/02/2009

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

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1325

Date of authorisation status change:
5/06/2023

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.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.