Salmoporc, lyophilisate and solvent for suspension for injection for pigs

Authorised

- Salmonella enterica, subsp. enterica, serovar Typhimurium, Live

Product identification

Medicine name:
Salmoporc, lyophilisate and solvent for suspension for injection for pigs
Salmoporc Lyofilisaat en oplosmiddel voor suspensie voor injectie
Salmoporc Lyophilisat et solvant pour suspension injectable
Salmoporc Lyophilisat und Lösungsmittel zur Herstellung einer Injektionssuspension

Active substance:
Salmonella enterica, subsp. enterica, serovar Typhimurium, Live

Target species:
Pig

Route of administration:
Oral use
Subcutaneous use

Product details

Active substance and strength:
Salmonella enterica, subsp. enterica, serovar Typhimurium, 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:
Belgium

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:
7/05/2019

Manufacturing sites for batch release:
Ceva-Phylaxia Veterinary Biologicals Co. Ltd.
IDT Biologika GmbH

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V541502

Date of authorisation status change:
7/05/2019

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)

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.

Package Leaflet

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