Salmoporc, lyophilisate for oral suspension for pigs

Authorised

- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 421/125 (histidine-adenine auxotrohic), Live

Product identification

Medicine name:
Salmoporc, lyophilisate for oral suspension for pigs
Salmoporc Lyofilisaat voor suspensie voor oraal gebruik
Salmoporc Lyophilisat pour suspension buvable
Salmoporc Lyophilisat zur Herstellung einer Suspension zum Einnehmen

Active substance:
Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 421/125 (histidine-adenine auxotrohic), Live

Target species:
Pig

Route of administration:
Oral use

Product details

Active substance and strength:

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 421/125 (histidine-adenine auxotrohic), Live
8.00 log10 colony forming unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral lyophilisate

Withdrawal period by route of administration:
Oral use:

- **Pig**
  - Meat and offal. 6 week

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 one 10 ml glass vial (type I) with rubber stoppers and aluminium crimp caps containing 200 doses (1 dose = 1 ml) of lyophilisate.

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 Zrt.
IDT Biologika GmbH

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V541511

Date of authorisation status change:
7/05/2019

Reference member state:
Netherlands

Procedure number:
NL/V/0247/002

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.

Labelling

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