Salmoporc, lyophilisate and solvent for suspension for injection / lyophilisate for use in drinking water

Authorised

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

Product identification

Medicine name:
Salmoporc, lyophilisate and solvent for suspension for injection / lyophilisate for use in drinking water
SALMOPORC LYOPHILISAT ET SOLVANT POUR SUSPENSION INJECTABLE / LYOPHILISAT POUR ADMINISTRATION DANS L'EAU DE BOISSON

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

Target species:
Pig (suckling piglet)
Pig (sow for reproduction)
Pig

Route of administration:

Oral use
Subcutaneous use

Product details

Active substance and strength:
Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 421/125 (histidine-adenine auxotrohic), Live
500000000.00 Colony forming unit / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:
Oral use:

- **Pig (suckling piglet)**
  - Meat and offal. 6 week

Subcutaneous use:

- **Pig (sow for reproduction)**
  - Meat and offal. 6 week

- **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:
France

Package description:
(ID2) 200 Dose: Box (Cardboard) with 1 Bottle (Glass) with 200 Dose
(ID1) 20 Dose; 20 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 20 millilitre(s) and 1 Bottle (Glass) with 20 Dose

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 Sante Animale

Marketing authorisation date:
16/08/2023

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.
IDT Biologika GmbH

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/7172249 4/2023

Date of authorisation status change:
16/08/2023

Reference member state:
Germany

Procedure number:

DE/V/0290/001

Concerned member states:
France Spain

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

Package Leaflet and Labelling

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