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 auxotrophic), Live

Product identification

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

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

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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.

Summary of Product Characteristics

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