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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

Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

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

Marketing authorisation date:
14/07/2021

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
4027 ESP

Date of authorisation status change:
15/07/2021

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

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.

Combined File of all Documents