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

Authorised

- Salmonella enterica, subsp. enterica, serovar Typhimurium, histidine-adenine auxotrophic, Live

Product identification

Medicine name:
Salmoporc, lyophilisate and solvent for suspension for injection / lyophilisate for use in drinking water
Salmoporc, Lyophilisat und Lösungsmittel zur Herstellung einer Suspension zur Injektion / Lyophilisat zur Anwendung mit Trinkwasser

Active substance:
Salmonella enterica, subsp. enterica, serovar Typhimurium, histidine-adenine auxotrophic, Live

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

Route of administration:
Subcutaneous use
Oral use

Product details

Active substance and strength:
Salmonella enterica, subsp. enterica, serovar Typhimurium, histidine-adenine auxotrophic, Live
5000.00 million colony forming units / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

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

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

Oral use:

- **Pig (suckling piglet)**
  - 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:
Germany

Package description:
(ID2): Cardboard box containing 1 vial with 200 doses
(ID1): Cardboard box containing 1 vial with 20 doses lyophilised vaccine and 1 vial with 20 ml solvent

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 Tiergesundheit GmbH

Marketing authorisation date:
25/07/2002

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

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.02340.01.1

Date of authorisation status change:
14/06/2007

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

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