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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

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Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

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 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

Documents

Combined File of all Documents

English (PDF)
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