BIOSUIS Salm emulsion for injection for pigs

Authorised

- Salmonella enterica, subsp. enterica, serovar Typhimurium, Inactivated
- Salmonella enterica, subsp. enterica, serovar Derby, Inactivated
- Salmonella enterica, subsp. enterica, serovar Infantis, Inactivated

Product identification

Medicine name:
BIOSUIS Salm emulsion for injection for pigs

Active substance:
Salmonella enterica, subsp. enterica, serovar Typhimurium, Inactivated
Salmonella enterica, subsp. enterica, serovar Derby, Inactivated
Salmonella enterica, subsp. enterica, serovar Infantis, Inactivated

Target species:
Pig (pregnant sow)
Pig (pregnant gilt)

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Salmonella enterica, subsp. enterica, serovar Typhimurium, Inactivated
1.00 relative potency / 1.00 Dose

Salmonella enterica, subsp. enterica, serovar Derby, Inactivated
1.00 relative potency / 1.00 Dose

Salmonella enterica, subsp. enterica, serovar Infantis, Inactivated
1.00 relative potency / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig (pregnant sow)**
  - Meat and offal. 0 day

- **Pig (pregnant gilt)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB14

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Netherlands

Package description:

Glass Vial 1 x 10.0 Dose

Plastic Vial 1 x 50.0 Dose
Plastic Vial 1 x 100.0 Dose
Glass Vial 10 x 10.0 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:
Bioveta a.s.

Marketing authorisation date:
8/11/2019

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 123500

Date of authorisation status change:
28/01/2022

Reference member state:
Czechia

Procedure number:
CZ/V/0151/001

Concerned member states:
Austria Belgium Estonia France Germany Greece Hungary Ireland Italy Netherlands Poland Slovakia United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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