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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:

Belgium

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:
26/11/2019

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
26/11/2019

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

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.

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