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Porcilis Glässer Vet Suspension for injection for pigs

Authorised

- Haemophilus parasuis, serotype 5, strain 4800, Inactivated
- Haemophilus parasuis, serotype 5, strain 4800, Inactivated
- Haemophilus parasuis, serotype 5, strain 4800, Inactivated

Product identification

Medicine name:
Porcilis Glässer Vet Suspension for injection for pigs

Active substance:
Haemophilus parasuis, serotype 5, strain 4800, Inactivated
Haemophilus parasuis, serotype 5, strain 4800, Inactivated
Haemophilus parasuis, serotype 5, strain 4800, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Haemophilus parasuis, serotype 5, strain 4800, Inactivated

9.10 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

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Haemophilus parasuis, serotype 5, strain 4800, Inactivated
9.10 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig**
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AB07

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Czechia

Available in:
Czechia

Package description:
6 x 20 ml (10 doses) in glass type I vials in carton (60 doses)
100 ml (50 doses) in PET vials in carton
6 x 50 ml (25 doses) in glass type I vials in carton (150 doses)
12 x 20 ml (10 doses) in PET vials in carton (120 doses)
20 ml (10 doses) in PET vials in carton

50 ml (25 doses) in glass type I vials in carton
100 ml (50 doses) in glass type I vials in carton
20 ml (10 doses) in glass type I vials in carton
6 x 20 ml (10 doses) in PET vials in carton (60 doses)
6 x 50 ml (25 doses) in PET vials in carton (150 doses)
12 x 100 ml (50 doses) in glass type I vials in carton (600 doses)
12 x 50 ml (25 doses) in PET vials in carton (300 doses)
12 x 50 ml (25 doses) in glass type I vials in cardboard box (300 doses)
6 x 100 ml (50 doses) in glass type I vials in carton (300 doses)
12 x 100 ml (50 doses) in PET vials in carton (600 doses)
12 x 20 ml (10 doses) in glass type I vials in carton (120 doses)
6 x 100 ml (50 doses) in PET vials in cardboard box (300 doses)
50 ml (25 doses) in PET vials in cardboard box

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
25/02/2004

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/020/04-C

Date of authorisation status change:

28/04/2008

Reference member state:
Denmark

Procedure number:
DK/V/0107/001

Concerned member states:
Austria Belgium Cyprus Czechia Estonia Finland France Germany Greece Hungary Ireland Italy Latvia Luxembourg Netherlands Norway Poland Portugal Slovakia Slovenia Spain 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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