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Porcilis Glässer Vet Suspension for injection for pigs

Not authorised

- 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

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)

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:

Surrendered

Authorised in:

Slovenia

Package description:

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

12 x 20 ml (10 doses) in glass type I vials in carton (120 doses)
6 x 50 ml (25 doses) in glass type I vials in carton (150 doses)

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:
9/12/2005

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

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
MR/V/0267/001

Date of authorisation status change:
31/08/2023

Reference member state:
Denmark

Procedure number:
DK/V/0107/001

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.

Labelling

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.

PI.pdf