File downloaded on 2026-11-14
**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000060320

# Porcilis Glässer

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

**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)
Haemophilus parasuis, serotype 5, strain 4800, Inactivated

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

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

**Authorisation status:**
Valid

**Authorised in:**
Ireland

**Available in:**
Ireland

**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 (Ireland) Limited

**Marketing authorisation date:**
4/03/2004

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

**Responsible authority:**
Health Products Regulatory Authority

**Authorisation number:**
VPA10996/179/001

**Date of authorisation status change:**
4/03/2004

**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

PI.pdf