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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:
Denmark

Available in:
Denmark

Package description:
Available only in Danish
Available only in Danish
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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:
13/03/2003

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

Responsible authority:
Danish Medicines Agency

Authorisation number:
31892

Date of authorisation status change:

13/03/2003

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

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