Porcilis PRRS, Lyofilizát a rozpouštědlo pro injekční suspenzi

Authorised

- Porcine reproductive and respiratory syndrome virus, type 1, strain DV, Live

Product identification

Medicine name:
Porcilis PRRS, Lyofilizát a rozpouštědlo pro injekční suspenzi

Active substance:
Porcine reproductive and respiratory syndrome virus, type 1, strain DV, Live

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Porcine reproductive and respiratory syndrome virus, type 1, strain DV, Live
6.30 log10 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AD03

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

Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech
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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:
Intervet International B.V.

Marketing authorisation date:
10/09/2001

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

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

Authorisation number:
97/036/01-C

Date of authorisation status change:
8/06/2011

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.