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ReproCyc PRRS EU lyophilisate and solvent for suspension for injection for pigs

Authorised

- Porcine reproductive and respiratory syndrome virus, type 1, strain PRRS 94881, Live

Product identification

Medicine name:
ReproCyc PRRS EU lyophilisate and solvent for suspension for injection for pigs

Active substance:
Porcine reproductive and respiratory syndrome virus, type 1, strain PRRS 94881, Live

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Porcine reproductive and respiratory syndrome virus, type 1, strain PRRS 94881, Live 10000000.00 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:
Germany

Available in:
Germany

Package description:
Lyophilisate:Type I amber glass vials with bromobutyl rubber stopper and aluminium seal.Solvent:High density polyethylene (HDPE) vials with a bromo- or chlorobutyl rubber stopper and aluminium seal. Cardboard box of 25 lyophilisate vials of 200 ml (100 doses) and Cardboard box of 25 solvent vials of 200 ml (100 doses).
Lyophilisate:Type I amber glass vials with bromobutyl rubber stopper and aluminium seal.Solvent:High density polyethylene (HDPE) vials with a bromo- or chlorobutyl rubber stopper and aluminium seal. Cardboard box of 25 lyophilisate vials of 100 ml (50 doses) and Cardboard box of 25 solvent vials of 100 ml (50 doses).
Lyophilisate:Type I amber glass vials with bromobutyl rubber stopper and aluminium seal.Solvent:High density polyethylene (HDPE) vials with a bromo- or chlorobutyl rubber stopper and aluminium seal. Cardboard box of 25 lyophilisate vials of 20 ml (10 doses) and Cardboard box of 25 solvent vials of 20 ml (10 doses).

Lyophilisate:Type I amber glass vials with bromobutyl rubber stopper and aluminium seal.Solvent:High density polyethylene (HDPE) vials with a bromo- or chlorobutyl rubber stopper and aluminium seal. Cardboard box of 12 lyophilisate vials of 200 ml (100 doses) and Cardboard box of 12 solvent vials of 200 ml (100 doses).
Lyophilisate:Type I amber glass vials with bromobutyl rubber stopper and aluminium seal.Solvent:High density polyethylene (HDPE) vials with a bromo- or chlorobutyl rubber stopper and aluminium seal. Cardboard box of 12 lyophilisate vials of 100 ml (50 doses) and Cardboard box of 12 solvent vials of 100 ml (50 doses).
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Lyophilisate:Type I amber glass vials with bromobutyl rubber stopper and aluminium seal.Solvent:High density polyethylene (HDPE) vials with a bromo- or chlorobutyl rubber stopper and aluminium seal. Cardboard box of 1 lyophilisate vial of 200 ml and 1 solvent vial of 200 ml (100 doses).
Lyophilisate:Type I amber glass vials with bromobutyl rubber stopper and aluminium seal.Solvent:High density polyethylene (HDPE) vials with a bromo- or chlorobutyl rubber stopper and aluminium seal. Cardboard box of 1 lyophilisate vial of 100 ml and 1 solvent vial of 100 ml (50 doses).
Lyophilisate:Type I amber glass vials with bromobutyl rubber stopper and aluminium seal.Solvent:High density polyethylene (HDPE) vials with a bromo- or chlorobutyl rubber stopper and aluminium seal. Cardboard box of 1 lyophilisate vial of 20 ml and 1 solvent vial of 20 ml (10 doses).

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:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

24/03/2015

Manufacturing sites for batch release:
Boehringer Ingelheim Vetmedica GmbH

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.11723.01.1

Date of authorisation status change:
20/04/2020

Reference member state:
Ireland

Procedure number:
IE/V/0444/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia France Germany Greece Hungary Italy Latvia Liechtenstein Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

English (PDF)

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Summary of Product Characteristics