Porcilis Begonia DF Suspension for intamuscular injection

Authorised

- Aujeszky's disease virus, strain Begonia gE- tk-, Live

Product identification

Medicine name:
Porcilis Begonia DF Suspension for intamuscular injection

Active substance:
Aujeszky's disease virus, strain Begonia gE- tk-, Live

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Aujeszky's disease virus, strain Begonia gE- tk-, Live
316228.00 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AD01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Luxembourg

Package description:

(ID12) 1000 Dose; 2000 millilitre(s): Box (Cardboard) with 10 Box (Cardboard) each with 1 Bottle (Glass type I) with 100 Dose, closed with Kappe and Stopfen (Aluminium, Halobutyl Rubber) and 1 Bottle (Glass) with 200 millilitre(s), closed with Kappe and Stopfen (Aluminium, Butylkautschuk)

(ID11) 500 Dose; 1000 millilitre(s): Box (Cardboard) with 5 Box (Cardboard) each with 1 Bottle (Glass type I) with 100 Dose, closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium) and 1 Bottle (Glass) with 200 millilitre(s), closed with Kappe and Stopfen (Aluminium, Butylkautschuk)

(ID10) 100 Dose; 200 millilitre(s): Box (Cardboard) with 1 Bottle (Glass type I) with 100 Dose, closed with Kappe and Stopfen (Aluminium, Halobutyl Rubber) and 1 Bottle (Glass) with 200 millilitre(s), closed with Kappe and Stopfen (Aluminium, Butylkautschuk)

(ID9) 500 Dose; 1000 millilitre(s): Box (Cardboard) with 10 Box (Cardboard) each with 1 Bottle (Glass type I) with 50 Dose, closed with Kappe and Stopfen (Aluminium, Halobutyl Rubber) and 1 Bottle (Glass) with 100 millilitre(s), closed with Kappe and Stopfen (Aluminium, Butylkautschuk)

(ID8) 250 Dose; 500 millilitre(s): Box (Cardboard) with 5 Box (Cardboard) each with 1 Bottle (Glass type I) with 50 Dose, closed with Kappe and Stopfen (Aluminium,

Halobutyl Rubber) and 1 Bottle (Glass) with 100 millilitre(s), closed with Kappe and Stopfen (Aluminium, Butylkautschuk)
(ID7) 50 Dose; 100 millilitre(s): Box (Cardboard) with 1 Bottle (Glass type I) with 50 Dose, closed with Kappe and Stopfen (Aluminium, Halobutyl Rubber) and 1 Bottle (Glass) with 100 millilitre(s), closed with Kappe and Stopfen (Aluminium, Butylkautschuk)
(ID6) 250 Dose; 500 millilitre(s): Box (Cardboard) with 10 Box (Cardboard) each with 1 Bottle (Glass type I) with 25 Dose, closed with Kappe and Stopfen (Aluminium, Halobutyl Rubber) and 1 Bottle (Glass) with 50 millilitre(s), closed with Kappe and Stopfen (Aluminium, Butylkautschuk)
(ID5) 125 Dose; 250 millilitre(s): Box (Cardboard) with 5 Box (Cardboard) each with 1 Bottle (Glass type I) with 25 Dose, closed with Kappe and Stopfen (Aluminium, Halobutyl Rubber) and 1 Bottle (Glass) with 50 millilitre(s), closed with Kappe and Stopfen (Aluminium, Butylkautschuk)
(ID4) 25 Dose; 50 millilitre(s): Box (Cardboard) with 1 Bottle (Glass type I) with 25 Dose, closed with Kappe and Stopfen (Aluminium, Halobutyl Rubber) and 1 Bottle (Glass) with 50 millilitre(s), closed with Kappe and Stopfen (Aluminium, Butylkautschuk)
(ID3) 100 Dose; 200 millilitre(s): Box (Cardboard) with 10 Box (Cardboard) each with 1 Bottle (Glass type I) with 10 Dose, closed with Kappe and Stopfen (Aluminium, Halobutyl Rubber) and 1 Bottle (Glass) with 20 millilitre(s), closed with Kappe and Stopfen (Aluminium, Butylkautschuk)
(ID2) 50 Dose; 100 millilitre(s): Box (Cardboard) with 5 Box (Cardboard) each with 1 Bottle (Glass type I) with 10 Dose, closed with Kappe and Stopfen (Aluminium, Halobutyl Rubber) and 1 Bottle (Glass) with 20 millilitre(s), closed with Kappe and Stopfen (Aluminium, Butylkautschuk)
(ID1) 10 Dose; 20 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 10 Dose, closed with Kappe and Stopfen (Aluminium, Halobutyl Rubber) and 1 Bottle (Glass) with 20 millilitre(s), closed with Kappe and Stopfen (Aluminium, Butylkautschuk)
(ID24) 1000 Dose; 2000 millilitre(s): Box (Cardboard) with 10 Box (Cardboard) each with 1 Bottle (Glass type I) with 100 Dose, closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium) and 1 Bottle (Polyethylenterephthalat) with 200 millilitre(s), closed with Stopfen and Kappe (Butylkautschuk, Aluminium)
(ID23) 500 Dose; 1000 millilitre(s): Box (Cardboard) with 5 Box (Cardboard) each with 1 Bottle (Glass type I) with 100 Dose, closed with Kappe and Stopfen (Aluminium, Halobutyl Rubber) and 1 Bottle (Polyethylenterephthalat) with 200 millilitre(s), closed with Stopfen and Kappe (Butylkautschuk, Aluminium)

(ID22) 100 Dose; 200 millilitre(s): Box (Cardboard) with 1 Bottle (Glass type I) with 100 Dose, closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium) and 1 Bottle (Polyethylenterephthalat) with 200 millilitre(s), closed with Stopfen and Kappe (Butylkautschuk, Aluminium)
(ID21) 500 Dose; 1000 millilitre(s): Box (Cardboard) with 10 Box (Cardboard) each with 1 Bottle (Glass type I) with 50 Dose, closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium) and 1 Bottle (Polyethylenterephthalat) with 100 millilitre(s), closed with Stopfen and Kappe (Butylkautschuk, Aluminium)
(ID20) 250 Dose; 500 millilitre(s): Box (Cardboard) with 5 Box (Cardboard) each with 1 Bottle (Glass type I) with 50 Dose, closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium) and 1 Bottle (Polyethylenterephthalat) with 100 millilitre(s), closed with Stopfen and Kappe (Butylkautschuk, Aluminium)
(ID19) 50 Dose; 100 millilitre(s): Box (Cardboard) with 1 Bottle (Glass type I) with 50 Dose, closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium) and 1 Bottle (Polyethylenterephthalat) with 100 millilitre(s), closed with Stopfen and Kappe (Butylkautschuk, Aluminium)
(ID18) 250 Dose; 500 millilitre(s): Box (Cardboard) with 10 Box (Cardboard) each with 1 Bottle (Glass type I) with 25 Dose, closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium) and 1 Bottle (Polyethylenterephthalat) with 50 millilitre(s), closed with Stopfen and Kappe (Butylkautschuk, Aluminium)
(ID17) 125 Dose; 250 millilitre(s): Box (Cardboard) with 5 Box (Cardboard) each with 1 Bottle (Glass type I) with 25 Dose, closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium) and 1 Bottle (Polyethylenterephthalat) with 50 millilitre(s), closed with Stopfen and Kappe (Butylkautschuk, Aluminium)
(ID16) 25 Dose; 50 millilitre(s): Box (Cardboard) with 1 Bottle (Glass type I) with 25 Dose, closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium) and 1 Bottle (Polyethylenterephthalat) with 50 millilitre(s), closed with Stopfen and Kappe (Butylkautschuk, Aluminium)
(ID15) 100 Dose; 200 millilitre(s): Box (Cardboard) with 10 Box (Cardboard) each with 1 Bottle (Glass type I) with 10 Dose, closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium) and 1 Bottle (Polyethylenterephthalat) with 20 millilitre(s), closed with Stopfen and Kappe (Butylkautschuk, Aluminium)
(ID14) 50 Dose; 100 millilitre(s): Box (Cardboard) with 5 Box (Cardboard) each with 1 Bottle (Glass type I) with 10 Dose, closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium) and 1 Bottle (Polyethylenterephthalat) with 20 millilitre(s), closed with Stopfen and Kappe (Butylkautschuk, Aluminium)

(ID13) 10 Dose; 20 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 10 Dose, closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium) and 1 Bottle (Polyethylenterephthalat) with 20 millilitre(s), closed with Stopfen and Kappe (Butylkautschuk, Aluminium)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
21/01/1993

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

Responsible authority:
Ministry Of Health And Social Security

Authorisation number:
V 817/96/12/0544

Date of authorisation status change:
21/01/1993

Reference member state:
Germany

Procedure number:
DE/V/0012/001

Concerned member states:

Belgium France Greece Italy Luxembourg Netherlands Portugal Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents