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Porcilis Begonia Unisolve Suspension for intramuscular injection

Authorised

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

Product identification

Medicine name:

Porcilis Begonia Unisolve Suspension for intramuscular 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 suspension 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

Authorisation status:

Valid

Authorised in:

Belgium

Package description:

(ID12) 500 Dose; 1000 millilitre(s): Box (cardboard) with 5 Bottle (Glass type I) each with 100 Dose, closed with Kappe (aluminium) and Stopper (Halobutyl Rubber) and 5 Bottle (Glass type II) each with 200 millilitre(s), closed with Kappe (aluminium) and Stopper (butyl rubber)

(ID10) 125 Dose; 250 millilitre(s): Box (cardboard) with 5 Bottle (Glass type I) each with 25 Dose, closed with Kappe (aluminium) and Stopper (Halobutyl Rubber) and 5 Bottle (Glass type I) each with 50 millilitre(s), closed with Kappe (aluminium) and Stopper (butyl rubber)

(ID11) 250 Dose; 500 millilitre(s): Box (cardboard) with 5 Bottle (Glass type I) each with 50 Dose, closed with Kappe (aluminium) and Stopper (Halobutyl Rubber) and 5 Bottle (Glass type II) each with 100 millilitre(s), closed with Kappe (aluminium) and Stopper (butyl rubber)

(ID9) 50 Dose; 100 millilitre(s): Box (cardboard) with 5 Bottle (Glass type I) each with 10 Dose, closed with Kappe (aluminium) and Stopper (Halobutyl Rubber) and 5 Bottle (Glass type I) each with 20 millilitre(s), closed with Kappe (aluminium) and Stopper (butyl rubber)

(ID1) 10 Dose; 20 millilitre(s): Box (cardboard) with 1 Bottle (Glass type I) with 10 Dose, closed with Stopper (Halobutyl Rubber) and Kappe (aluminium) and 1 Bottle (Glass type I) with 20 millilitre(s), closed with Stopper (butyl rubber) and Kappe (aluminium)

(ID2) 100 Dose; 200 millilitre(s): Box (cardboard) with 10 Bottle (Glass type I) each with 10 Dose, closed with Stopper (Halobutyl Rubber) and Kappe (aluminium) and 10 Bottle (Glass type I) each with 20 millilitre(s), closed with Stopper (butyl rubber) and Kappe (aluminium)

(ID3) 25 Dose; 50 millilitre(s): Box (cardboard) with 1 Bottle (Glass type I) with 25 Dose, closed with Stopper (Halobutyl Rubber) and Kappe (aluminium) and 1 Bottle (Glass type I) with 50 millilitre(s), closed with Stopper (butyl rubber) and Kappe (aluminium)

(ID4) 250 Dose; 500 millilitre(s): Box (cardboard) with 10 Bottle (Glass type I) each with 25 Dose, closed with Stopper (Halobutyl Rubber) and Kappe (aluminium) and 10 Bottle (Glass type I) each with 50 millilitre(s), closed with Stopper (butyl rubber) and Kappe (aluminium)

(ID5) 50 Dose; 100 millilitre(s): Box (cardboard) with 1 Bottle (Glass type I) with 50 Dose, closed with Stopper (Halobutyl Rubber) and Kappe (aluminium) and 1 Bottle (Glass type II) with 100 millilitre(s), closed with Stopper (butyl rubber) and Kappe (aluminium)

(ID6) 500 Dose; 1000 millilitre(s): Box (cardboard) with 10 Bottle (Glass type II) each with 100 millilitre(s), closed with Stopper (butyl rubber) and Kappe (aluminium) and 10 Bottle (Glass type I) each with 50 Dose, closed with Stopper (Halobutyl Rubber) and Kappe (aluminium)

(ID7) 100 Dose; 200 millilitre(s): Box (cardboard) with 1 Bottle (Glass type I) with 100 Dose, closed with Stopper (Halobutyl Rubber) and Kappe (aluminium) and 1 Bottle (Glass type II) with 200 millilitre(s), closed with Stopper (butyl rubber) and Kappe (aluminium)

(ID8) 1000 Dose; 2000 millilitre(s): Box (cardboard) with 10 Bottle (Glass type I) each with 100 Dose, closed with Stopper (Halobutyl Rubber) and Kappe (aluminium) and 10 Bottle (Glass type II) each with 200 millilitre(s), closed with Stopper (butyl rubber) and Kappe (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:

22/10/1991

Manufacturing sites for batch release:

Intervet International B.V.
Intervet International GmbH

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

7/08/2023

Reference member state:

Germany

Procedure number:

DE/V/0006/001

Concerned member states:

Belgium France Italy Netherlands Portugal

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.

Combined File of all Documents