Porcilis Begonia IDAL Suspension for intradermal injection

Authorised

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

Product identification

Medicine name:
Porcilis Begonia IDAL liofilizado e solvente para suspensão para injeção intradérmica em suínos
Porcilis Begonia IDAL Suspension for intradermal injection

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

Target species:
Pig

Route of administration:
Intradermal use

Product details

Active substance and strength:
Aujeszky's disease virus, strain Begonia gE- tk-, Live
3162280.00 tissue culture infective dose 50 / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Intradermal 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:

Portugal

Package description:

(ID15): 1 Box with 10 Box with (1 Bottle (Glass) with 100 Dose and 1 Bottle (PolyEthylene TerePhthalate) with 20 millilitre(s)) (1000.0 Dose, 200.0 millilitre(s))
(ID14): 1 Box with 10 Box with (1 Bottle (Glass) with 100 Dose and 1 Bottle (Glass) with 20 millilitre(s)) (1000.0 Dose, 200.0 millilitre(s))
(ID13): 1 Box with 5 Box with (1 Bottle (Glass) with 100 Dose and 1 Bottle (PolyEthylene TerePhthalate) with 20 millilitre(s)) (500.0 Dose, 100.0 millilitre(s))
(ID12): 1 Box with 5 Box with (1 Bottle (Glass) with 100 Dose and 1 Bottle (Glass) with 20 millilitre(s)) (500.0 Dose, 100.0 millilitre(s))
(ID11): 1 Box with (1 Bottle (Glass) with 100 Dose and 1 Bottle (PolyEthylene TerePhthalate) with 20 millilitre(s)) (100.0 Dose, 20.0 millilitre(s))
(ID10): 1 Box with (1 Bottle (Glass) with 100 Dose and 1 Bottle (Glass) with 20 millilitre(s)) (100.0 Dose, 20.0 millilitre(s))
(ID9): 1 Box with 10 Box with (1 Bottle (Glass) with 50 Dose and 1 Bottle (Glass) with 10 millilitre(s)) (500.0 Dose, 100.0 millilitre(s))
(ID8): 1 Box with 5 Box with (1 Bottle (Glass) with 10 millilitre(s) and 1 Bottle (Glass) with 50 Dose) (50.0 millilitre(s), 250.0 Dose)
(ID7): 1 Box with (1 Bottle (Glass) with 50 Dose and 1 Bottle (Glass) with 10 millilitre(s)) (50.0 Dose, 10.0 millilitre(s))
(ID6): 1 Box with 10 Box with (1 Bottle (Glass) with 25 Dose and 1 Bottle (Glass) with 5 millilitre(s)) (250.0 Dose, 50.0 millilitre(s))

(ID5): 1 Box with 5 Box with (1 Bottle (Glass) with 25 Dose and 1 Bottle (Glass) with 5 millilitre(s)) (125.0 Dose, 25.0 millilitre(s))
(ID4): 1 Box with (1 Bottle (Glass) with 5 millilitre(s) and 1 Bottle (Glass) with 25 Dose) (5.0 millilitre(s), 25.0 Dose)
(ID3): 1 Box with 10 Box with (1 Bottle (Glass) with 10 Dose and 1 Bottle (Glass) with 2 millilitre(s)) (100.0 Dose, 20.0 millilitre(s))
(ID2): 1 Box with 5 Box with (1 Bottle (Glass) with 10 Dose and 1 Bottle (Glass) with 2 millilitre(s)) (50.0 Dose, 10.0 millilitre(s))
(ID1): 1 Box with (1 Bottle (Glass) with 10 Dose and 1 Bottle (Glass) with 2 millilitre(s)) (10.0 Dose, 2.0 millilitre(s))

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:
MSD Animal Health Lda.

Marketing authorisation date:
2/02/1996

Manufacturing sites for batch release:
INTERVET INTERNATIONAL B.V.

Responsible authority:
DGAV

Authorisation number:
546/95 DGV

Date of authorisation status change:
2/02/1996

Reference member state:
Germany

Procedure number:
DE/V/0019/001

Concerned member states:
Belgium France Greece Italy Netherlands Portugal 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