AD live-SUIVAX

Authorised

- Aujeszky's disease virus, strain LomBart gE-, Live

Product identification

Medicine name:
AD live-SUIVAX

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

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Aujeszky's disease virus, strain LomBart gE-, Live
6.50 log10 50% cell culture infectious dose / 2.00 millilitre(s)

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:
Portugal

Package description:
Cardboard box containing 10 glass vials of 50 doses of vaccine sealed with rubber cap and aluminum ring with rubber cap and aluminum ring + 10 glass or polypropylene vials of 100 ml of glass or polypropylene bottles of 100 ml of diluent with elastomer cap and aluminum ring.
Cardboard box containing 10 glass vials of 10 doses of vaccine sealed with rubber stopper and aluminum ring + 10 glass or polypropylene vials of 20 ml of diluent with elastomer stopper and aluminum ring.
Cardboard box containing 50 doses glass vial of sealed vaccine with rubber stopper and aluminum ring + 100 ml vial or polypropylene bottle of diluent with elastomer stopper and aluminum ring.
Cardboard box containing 10 doses glass vial of sealed vaccine with rubber stopper and aluminum ring + 1 20 ml glass or polypropylene vial of diluent with elastomer stopper and aluminum ring.

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:
Fatro S.p.A.

Marketing authorisation date:
24/04/2002

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:

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Date of authorisation status change:
24/04/2002

Reference member state:
Italy

Procedure number:
IT/V/0109/001

Concerned member states:
Portugal Spain

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

Documents

Combined File of all Documents

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