AD-LIVE-SUIVAX, κόνις και διαλύτης για ενέσιμο εναιώρημα

Authorised

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

Product identification

Medicine name:
AD-LIVE-SUIVAX, κόνις και διαλύτης για ενέσιμο εναιώρημα

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
5.50 log10 50% tissue culture infectious dose / 2.00 millilitre(s)

Pharmaceutical form:
Powder and solvent for suspension for injection

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

Package description:
Available only in Greek
Available only in Greek

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
4/11/1999

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

Responsible authority:
National Organization For Medicines

Authorisation number:
37646/05-11-1999/K-0126701

Date of authorisation status change:
5/10/2016

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