OVALAXIA ενέσιμο εναιώρημα για πρόβατα

Authorised

- Mycoplasma agalactiae, strain AG 8, Inactivated

Product identification

Medicine name:
OVALAXIA ενέσιμο εναιώρημα για πρόβατα

Active substance:
Mycoplasma agalactiae, strain AG 8, Inactivated

Target species:
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Mycoplasma agalactiae, strain AG 8, Inactivated
9.00 log10 colour changing units / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI04AB

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:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
29/05/2019

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

Responsible authority:
National Organization For Medicines

Authorisation number:
15840/13-02-2025/K-0213301

Date of authorisation status change:
15/03/2020

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.