OVAX CLAMIDIA, ενέσιμο εναιώρημα για πρόβατα

Authorised

- Chlamydia abortus, Inactivated

Product identification

Medicine name:
OVAX CLAMIDIA, ενέσιμο εναιώρημα για πρόβατα

Active substance:
Chlamydia abortus, Inactivated

Target species:
Sheep

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Chlamydia abortus, Inactivated
5.00 log 10 50% embryo lethal dose / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Subcutaneous use:

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

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:
7/10/1999

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

Responsible authority:
National Organization For Medicines

Authorisation number:
130059/30-11-2022/K-0123801

Date of authorisation status change:
29/11/2022

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.