

COGLAREV ΚΟΝΙΣ ΚΑΙ ΔΙΑΛΥΤΗΣ ΓΙΑ ΕΝΑΙΩΡΗΜΑ

Authorised

- Brucella melitensis, strain REV 1, Live

Product identification

Medicine name:

COGLAREV ΚΟΝΙΣ ΚΑΙ ΔΙΑΛΥΤΗΣ ΓΙΑ ΕΝΑΙΩΡΗΜΑ

Active substance:

Brucella melitensis, strain REV 1, Live

Target species:

Sheep (lamb)

Goat (kid)

Route of administration:

Intraocular use

Product details

Active substance and strength:

Brucella melitensis, strain REV 1, Live

1000000000.00 plaque forming unit / 1.00 Dose

Pharmaceutical form:

Powder and solvent for intraocular instillation solution

Withdrawal period by route of administration:

Intraocular use:**• Sheep (lamb)**

- Meat and offal. 3 month

και σύμφωνα με την ισχύουσα νομοθεσία για τη βρουκέλλωση

• Goat (kid)

- Meat and offal. 3 month

και σύμφωνα με την ισχύουσα νομοθεσία για τη βρουκέλλωση

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Package description:

Available only in Greek

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone)

Marketing authorisation holder:

Ceva Sante Animale

Marketing authorisation date:

9/02/2004

Manufacturing sites for batch release:

Cz Veterinaria S.A.

Responsible authority:

National Organization For Medicines

Authorisation number:

7713/09-02-2004/K-0147101

Date of authorisation status change:

8/02/2017

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000982603>