OCUREV

Authorised

- Brucella melitensis, strain REV 1, Live

Product identification

Medicine name:
OCUREV
OCUREV

Active substance:
Brucella melitensis, strain REV 1, Live

Target species:
Sheep
Goat

Route of administration:
Ocular use

Product details

Active substance and strength:
Brucella melitensis, strain REV 1, Live
2000000000.00 Colony forming unit / 1.00 Dose

Pharmaceutical form:
Eye drops, powder and solvent for suspension

Withdrawal period by route of administration:
Ocular use:

- **Sheep**
  - Meat and offal. 30 day
- **Goat**
  - Meat and offal. 30 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI04AE

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

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
1 box containing 1 vial of 25 doses and 1 vial of 1 ml of solvent
1 box containing 1 vial of 50 doses and 1 vial of 2 ml of solvent
1 box containing 1 vial of 10 doses and 1 vial of 0.5 ml of solvent

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
CZ Vaccines S.A.U.

Marketing authorisation date:
6/07/2017

Manufacturing sites for batch release:
Cz Veterinaria S.A.

Responsible authority:
National Veterinary Medicines Agency

Authorisation number:
FR/V/1723018 6/2017

Date of authorisation status change:
6/07/2017

Reference member state:
Spain

Procedure number:
ES/V/0107/001

Concerned member states:
France Portugal

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 21/12/2023
Download

Combined File of all Documents

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

Labelling
