OCUREV eye drops, powder and solvent for suspension

Authorised

- Brucella melitensis, strain REV 1, Live

Product identification

Medicine name:
OCUREV eye drops, powder and solvent for suspension

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:

Portugal

Available in:

Portugal

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:
20/11/2002

Manufacturing sites for batch release:
CZ Vaccines S.A.U.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
R744/05DGV

Date of authorisation status change:
26/08/2022

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

Combined File of all Documents

English (PDF)

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Package Leaflet

Labelling

Summary of Product Characteristics