

OVILIS TOXOVAX CONCENTRATE AND SOLVENT FOR SUSPENSION FOR INJECTION FOR SHEEP

Authorised

- Toxoplasma gondii, strain S48, Live

Product identification

Medicine name:

OVILIS TOXOVAX CONCENTRATE AND SOLVENT FOR SUSPENSION FOR INJECTION FOR SHEEP

Active substance:

Toxoplasma gondii, strain S48, Live

Target species:

Sheep

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Toxoplasma gondii, strain S48, Live
100000.00 Organisms / 1.00 Dose

Pharmaceutical form:

Concentrate and solvent for suspension for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Sheep

- Meat and offal. 42 day
 - Milk. 0 day
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI04AN01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Package description:

Carton box with 1 vial of 20 doses of vaccine and 1 vial of 40 ml diluents

1 vial of 100 ml of diluent

1 vial of 40 ml of diluent

1 vial 50 doses of vaccine

1 vial 20 doses of vaccine

Carton box with 1 vial of 50 doses of vaccine and 1 vial of 100 ml diluents

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Intervet Nederland B.V.

Marketing authorisation date:

19/10/2010

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 9827

Date of authorisation status change:

21/06/2022

Reference member state:

France

Procedure number:

FR/V/0119/001

Concerned member states:

Netherlands Spain

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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