

Toxovax

Authorised

- Toxoplasma gondii, strain S48, Live

Product identification

Medicine name:

Toxovax

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
7500000.00 other / 1.00 Dose

Pharmaceutical form:

Concentrate and solvent for suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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Sheep

- Meat and offal. 42 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI04AN01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

Carton box with 1 vial of 20 doses of vaccine and a vial of 40 ml of diluent, and a transfer spike.Vaccine:Type I Ph.Eur. glass vials, closed with a halogenobutyl rubber stopper and sealed with a colour coded aluminium cap,containing 2 ml (20 doses) of vaccine concentrate.Diluent:Hydrolytic Type II Ph.Eur. glass vials with halogenated butyl rubber stopper and an aluminium crimp cap containing 40 ml (for 20 doses) diluent

Carton box with 1 vial of 20 doses of vaccine and a vial of 40 ml of diluent, and a transfer spike.Vaccine:Type I Ph.Eur. glass vials, closed with a halogenobutyl rubber stopper and sealed with a colour coded aluminium cap,containing 2 ml (20 doses) of vaccine concentrate.Diluent:PET (polyethylene terephthalate) vials with halogenated butyl rubber stopper and an aluminium crimp cap containing 40 ml (for 20 doses) diluent

Carton box with 1 vial of 50 doses of vaccine and a carton box with 100ml of diluent (add transfer spike if appropriate to either carton). Vaccine:Type I Ph.Eur. glass vials, closed with a halogenobutyl rubber stopper and sealed with a colour coded aluminium cap, containing 5 ml (50 doses) of vaccine concentrate.Diluent:Hydrolytic Type II Ph. Eur. glass vials with halogenated butyl rubber stopper and an aluminium

crimp cap containing 100 ml (for 50 doses) diluent.

Carton box with 1 vial of 50 doses of vaccine and a carton box with 100ml of diluent (add transfer spike if appropriate to either carton). Vaccine:Type I Ph.Eur. glass vials, closed with a halogenobutyl rubber stopper and sealed with a colour coded aluminium cap, containing 5 ml (50 doses) of vaccine concentrate.Diluent:PET (polyethylene terephthalate) vials with halogenated butyl rubber stopper and an aluminium crimp cap containing 100 ml (for 50 doses) diluent.

Carton box with 1 vial of 50 doses of vaccine and a vial of 100ml of diluent, and a transfer spike.Vaccine:Type I Ph.Eur. glass vials, closed with a halogenobutyl rubber stopper and sealed with a colour coded aluminium cap, containing 5 ml (50 doses) of vaccine concentrate.Diluent:PET (polyethylene terephthalate) vials with halogenated butyl rubber stopper and an aluminium crimp cap containing 100 ml (for 50 doses) diluent.

Carton box with 1 vial of 20 doses of vaccine and a carton box with 40ml of diluent (add transfer spike if appropriate to either carton). Vaccine:Type I Ph.Eur. glass vials, closed with a halogenobutyl rubber stopper and sealed with a colour coded aluminium cap, containing 2 ml (20 doses) of vaccine concentrate.Diluent:Hydrolytic Type II Ph.Eur. glass vials with halogenated butyl rubber stopper and an aluminium crimp cap containing 20 ml (for 20 doses) diluent.

Carton box with 1 vial of 20 doses of vaccine and a carton box with 40ml of diluent (add transfer spike if appropriate to either carton). Vaccine:Type I Ph.Eur. glass vials, closed with a halogenobutyl rubber stopper and sealed with a colour coded aluminium cap, containing 2 ml (20 doses) of vaccine concentrate.Diluent:PET (polyethylene terephthalate) vials with halogenated butyl rubber stopper and an aluminium crimp cap containing 20 ml (for 20 doses) diluent.

Carton box with 1 vial of 50 doses of vaccine and a vial of 100ml of diluent, and a transfer spike.Vaccine:Type I Ph.Eur. glass vials, closed with a halogenobutyl rubber stopper and sealed with a colour coded aluminium cap, containing 5 ml (50 doses) of vaccine concentrate.Diluent:Hydrolytic Type II Ph.Eur. glass vials with halogenated butyl rubber stopper and an aluminium crimp cap containing 100 ml (for 50 doses) diluent.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet (Ireland) Limited

Marketing authorisation date:

18/08/2003

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10996/080/001

Date of authorisation status change:

18/08/2003

To consult adverse reactions on veterinary medicinal products please go to

www.adrreports.eu/vet

Documents

Summary of Product Characteristics