CEVAC TRANSMUNE lyophilisate for suspension for injection with solvent for chickens

Authorised

- Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Inactivated

Product identification

Medicine name:
CEVAC TRANSMUNE lyophilisate for suspension for injection with solvent for chickens
CEVAC TRANSMUNE liofilizzato con diluente per sospensione iniettabile per polli

Active substance:
Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Inactivated

Target species:
Chicken (embryonated eggs)
Chicken

Route of administration:
In ovo
Subcutaneous use

Product details

Active substance and strength:
Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Inactivated
0.10 50% Protective Dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD09

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

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Cardboard box with 20 glass vials (Type I) of 13.5 ml containing 8000 doses, closed with a bromobutyl stopper and sealed with aluminium caps with plastic tear-off centres.
Cardboard box with 20 glass vials (Type I) of 10 ml containing 5000 doses, closed with a bromobutyl stopper and sealed with aluminium caps with plastic tear-off centres.
Cardboard box with 20 glass vials (Type I) of 10 ml containing 4000 doses, closed with a bromobutyl stopper and sealed with aluminium caps with plastic tear-off centres.
Cardboard box with 20 glass vials (Type I) of 10 ml containing 2500 doses, closed with a bromobutyl stopper and sealed with aluminium caps with plastic tear-off centres.
Cardboard box with 20 glass vials (Type I) of 10 ml containing 2000 doses, closed with a bromobutyl stopper and sealed with aluminium caps with plastic tear-off centres.

Cardboard box with single glass vial (Type I) of 13.5 ml containing 8000 doses, closed with a bromobutyl stopper and sealed with aluminium caps with plastic tear-off centres.
Cardboard box with single glass vial (Type I) of 10 ml containing 5000 doses, closed with a bromobutyl stopper and sealed with aluminium caps with plastic tear-off centres.
Cardboard box with single glass vial (Type I) of 10 ml containing 4000 doses, closed with a bromobutyl stopper and sealed with aluminium caps with plastic tear-off centres.
Cardboard box with single glass vial (Type I) of 10 ml containing 2500 doses, closed with a bromobutyl stopper and sealed with aluminium caps with plastic tear-off centres.
Cardboard box with single glass vial (Type I) of 10 ml containing 2000 doses, closed with a bromobutyl stopper and sealed with aluminium caps with plastic tear-off centres.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Salute Animale S.p.A.

Marketing authorisation date:
13/02/2008

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.

Responsible authority:
Ministry Of Health

Authorisation number:

103966

Date of authorisation status change:
25/07/2012

Reference member state:
Hungary

Procedure number:
HU/V/0141/002

Concerned member states:
Bulgaria Cyprus Czechia Estonia France Germany Greece Ireland Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
Published on: 26/08/2025
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