HIPRAVIAR-TRT 4, Injekční emulze

Authorised

- Newcastle disease virus, strain La Sota, Inactivated
- Infectious bronchitis virus, strain H52, Inactivated
- Turkey rhinotracheitis virus, strain 1062, Inactivated
- Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Inactivated

Product identification

Medicine name:
HIPRAVIAR-TRT 4, Injekční emulze

Active substance:
Newcastle disease virus, strain La Sota, Inactivated

Infectious bronchitis virus, strain H52, Inactivated

Turkey rhinotracheitis virus, strain 1062, Inactivated

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

Target species:
Turkey
Chicken

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Newcastle disease virus, strain La Sota, Inactivated
16.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Infectious bronchitis virus, strain H52, Inactivated
288.00 enzyme-linked immunosorbent assay unit / 1.00 Dose

Turkey rhinotracheitis virus, strain 1062, Inactivated
195.00 enzyme-linked immunosorbent assay unit / 1.00 Dose

Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Inactivated
329.00 enzyme-linked immunosorbent assay unit / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Turkey**
  - Meat and offal. 0 day
  - Egg. 0 day
- **Chicken**
  - Meat and offal. 0 day
  - Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:
Czechia

Package description:
Available only in Czech

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:
Laboratorios Hipra S.A.

Marketing authorisation date:
15/04/2004

Manufacturing sites for batch release:
Laboratorios Hipra S.A.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/073/04-C

Date of authorisation status change:
8/06/2010

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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