HIPRAVIAR-TRT4

Authorised

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

Product identification

Medicine name:
HIPRAVIAR-TRT4

Active substance:
Turkey rhinotracheitis virus, strain 1062, Inactivated

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

Avian infectious bronchitis virus, strain H52, Inactivated

Newcastle disease virus, strain La Sota, Inactivated

Target species:
Chicken

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Turkey rhinotracheitis virus, strain 1062, Inactivated
6.00 log10 (50% tissue culture infectious dose)/dose / 1.00 Dose

Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Live
5.50 log 10 50% embryo infective dose / 1.00 Dose

Avian infectious bronchitis virus, strain H52, Inactivated
6.00 log 10 50% embryo infective dose / 1.00 Dose

Newcastle disease virus, strain La Sota, Inactivated
819.00 haemagglutinating units / 1.00 Dose

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AA06

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian

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:
Laboratorios Hipra S.A.

Marketing authorisation date:
9/03/2010

Manufacturing sites for batch release:
LABORATORIOS HIPRA, S.A.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2434/09.12.2014

Date of authorisation status change:
8/12/2014

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

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