

HIPRAVIAR-TRT4

Not
authorised

- Turkey rhinotracheitis virus, strain 1062, Inactivated
- Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Live
- 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

Infectious bronchitis virus, strain H52, Inactivated

Newcastle disease virus, strain La Sota, Inactivated

Target species:

Chicken (chick)

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Turkey rhinotracheitis virus, strain 1062, Inactivated

195.00 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

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

329.00 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Infectious bronchitis virus, strain H52, Inactivated

228.00 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Newcastle disease virus, strain La Sota, Inactivated

16.00 haemagglutination inhibiting unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Chicken (chick)

- 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:

Surrendered

Authorised in:

Romania

Package description:

Available only in Romanian

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:

19/04/2005

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

110037

Date of authorisation status change:

12/03/2025

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.