

# HIPRAVIAR-TRT4 EMULSION INYECTABLE PARA GALLINAS

Authorised

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

## Product identification

**Medicine name:**

HIPRAVIAR-TRT4 EMULSION INYECTABLE PARA GALLINAS

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**Active substance:**

Turkey rhinotracheitis virus, strain 1062, Inactivated

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

Newcastle disease virus, strain La Sota, Inactivated

Infectious bronchitis virus, strain H52, Inactivated

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**Target species:**

Chicken (pullet)

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**Route of administration:**

Intramuscular use

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## Product details

### **Active substance and strength:**

Turkey rhinotracheitis virus, strain 1062, Inactivated

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

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

357.00 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)

Infectious bronchitis virus, strain H52, Inactivated

2.40 serum neutralising unit(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Emulsion for injection

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### **Withdrawal period by route of administration:**

#### **Intramuscular use:**

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#### **Chicken (pullet)**

- Meat and offal. 0 day

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### **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AA06

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### **Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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### **Authorisation status:**

Valid

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### **Authorised in:**

Spain

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### **Package description:**

Available only in Spanish

Available only in [Spanish](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Laboratorios Hipra S.A.

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**Marketing authorisation date:**

3/07/1991

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**Manufacturing sites for batch release:**

Laboratorios Hipra S.A.

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

2373 ESP

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**Date of authorisation status change:**

6/10/2011

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Summary of Product Characteristics

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### Labelling

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.