BRONIPRA-ND/IBD

Authorised

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

Product identification

Medicine name:
BRONIPRA-ND/IBD

Active substance:
Avian infectious bronchitis virus, strain H52, Inactivated

Newcastle disease virus, strain La Sota, Inactivated

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

Target species:
Chicken (layer hen)
Chicken (for reproduction)

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Avian infectious bronchitis virus, strain H52, Inactivated
16.00 serum neutralising unit(s) / 1.00 Dose

Newcastle disease virus, strain La Sota, Inactivated
1024.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Inactivated
13500.00 enzyme-linked immunosorbent assay unit / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken (layer hen)**
  - Meat and offal. 0 day

- **Chicken (for reproduction)**
  - Meat and offal. 0 day

Subcutaneous use:

- **Chicken (layer hen)**
  - Meat and offal. 0 day

- **Chicken (for reproduction)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA08

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

Authorisation status:
Valid

Authorised in:
Spain

Package description:
Available only in Spanish
Available only in Spanish

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:
16/09/1988

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

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:
2297 ESP

Date of authorisation status change:
17/05/2011

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.