Gumbohatch (--) - Lyophilisate and solvent for suspension for injection

Authorised

- Infectious bursal disease virus, strain 1052 (intermediate plus), Live

Product identification

Medicine name:
Gumbohatch (--) - Lyophilisate and solvent for suspension for injection

Active substance:
Infectious bursal disease virus, strain 1052 (intermediate plus), Live

Target species:
Chicken
Chicken (embryonated eggs)

Route of administration:
In ovo
Subcutaneous use

Product details

Active substance and strength:
Infectious bursal disease virus, strain 1052 (intermediate plus), Live

Presentation_strength:$10^{1.48}$ - $10^{2.63}$ PU Reference:Hse Index:0

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

In ovo:

- **Chicken**
  - Not applicable. 0 day — Zero days
- **Chicken (embryonated eggs)**
  - Not applicable. 0 day — Zero days

Subcutaneous use:

- **Chicken**
  - Not applicable. 0 day — Zero days
- **Chicken (embryonated eggs)**
  - Not applicable. 0 day — Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania ,

Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:
Packaging:Vial (glass), Package_size:10 vials, Content:2500 doses
Packaging:Vial (glass), Package_size:10 vials, Content:2000 doses
Packaging:Vial (glass), Package_size:10 vials, Content:4000 doses
Packaging:Vial (glass), Package_size:10 vials, Content:5000 doses
Packaging:Vial (glass), Package_size:10 vials, Content:1000 doses
Packaging:Vial (glass), Package_size:10 vials, Content:10000 doses
Packaging:Vial (glass), Package_size:10 vials, Content:8000 doses

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:
12/11/2019

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

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
15/07/2022

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

Documents

Combined File of all Documents

English (PDF)
Published on: 25/11/2025
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