Vaxxinact H5 (--) - Emulsion for injection

Authorised

- Influenza A virus, subtype H5 (avian), haemagglutinin

Product identification

Medicine name:
Vaxxinact H5 (--) - Emulsion for injection

Active substance:
Influenza A virus, subtype H5 (avian), haemagglutinin

Target species:
Turkey
Chicken
Duck

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Influenza A virus, subtype H5 (avian), haemagglutinin

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Turkey**
  - All relevant tissues. 0 day Zero days
- **Chicken**
  - All relevant tissues. 0 day Zero days
- **Duck**
  - All relevant tissues. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA23

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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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:Bottle (HDPE), Package_size:20 bottles, Content:Bottle of 500 ml (1000 doses)

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:Bottle of 500 ml (1000 doses)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
New active substance - Exceptional circumstances (Article 25 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
4/12/2025

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

Responsible authority:
European Commission

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

Date of authorisation status change:
4/12/2025

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

Documents

Combined File of all Documents

English (PDF)
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