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PA-OLVAC +I+E Vaccino inattivato in emulsione iniettabile per tacchini

Authorised

- Turkey haemorrhagic enteritis virus, Inactivated
- Newcastle disease virus, Inactivated
- Riemerella anatipestifer, serotype 1, Inactivated
- Riemerella anatipestifer, serotype 3, Inactivated
- Influenza A virus, subtype H9N2 (avian), Inactivated
- Influenza A virus, subtype H6N2 (avian), Inactivated

Product identification

Medicine name:
PA-OLVAC +I+E Vaccino inattivato in emulsione iniettabile per tacchini

Active substance:
Turkey haemorrhagic enteritis virus, Inactivated

Newcastle disease virus, Inactivated

Riemerella anatipestifer, serotype 1, Inactivated

Riemerella anatipestifer, serotype 3, Inactivated

Influenza A virus, subtype H9N2 (avian), Inactivated

Influenza A virus, subtype H6N2 (avian), Inactivated

Target species:
Turkey

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Turkey haemorrhagic enteritis virus, Inactivated
1733.00 enzyme-linked immunosorbent assay unit / 0.50 millilitre(s)

Newcastle disease virus, Inactivated
50.00 50% Protective Dose / 0.50 millilitre(s)

Riemerella anatipestifer, serotype 1, Inactivated
1.00 billion colony forming units / 0.50 millilitre(s)

Riemerella anatipestifer, serotype 3, Inactivated
1.00 billion colony forming units / 0.50 millilitre(s)

Influenza A virus, subtype H9N2 (avian), Inactivated

Influenza A virus, subtype H6N2 (avian), Inactivated

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Turkey**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01CL01

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

Authorisation status:
Valid

Authorised in:

Italy

Package description:
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian

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:
Fatro S.p.A.

Marketing authorisation date:
9/03/2004

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Ministry Of Health

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

Date of authorisation status change:
9/03/2009

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

Documents

Combined File of all Documents

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