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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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