PHARMAVAC PHA emulsion for injection for pigeons

Authorised

- Pigeon paramyxovirus 1, strain 988M, Inactivated
- Pigeon herpesvirus, strain V298/70, Inactivated
- Fowl aviadenovirus 8, strain M2/E, Inactivated

Product identification

Medicine name:
PHARMAVAC PHA emulsion for injection for pigeons

Active substance:
Pigeon paramyxovirus 1, strain 988M, Inactivated

Pigeon herpesvirus, strain V298/70, Inactivated

Fowl aviadenovirus 8, strain M2/E, Inactivated

Target species:
Pigeon

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Pigeon paramyxovirus 1, strain 988M, Inactivated

6.90 log2 haemagglutination inhibiting unit(s) / 0.30 millilitre(s)

Pigeon herpesvirus, strain V298/70, Inactivated
38.10 enzyme-linked immunosorbent assay unit / 0.30 millilitre(s)

Fowl aviadenovirus 8, strain M2/E, Inactivated
24.70 enzyme-linked immunosorbent assay unit / 0.30 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Pigeon**
  - All relevant tissues. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01EA

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

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
Paper carton containing one glass vial, type I closed with chlorobutyl rubber stopper sealed with aluminium cap

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:
Pharmagal Bio spol. s r.o.

Marketing authorisation date:
26/10/2018

Manufacturing sites for batch release:
Pharmagal Bio spol. s r.o.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/055/DC/18

Date of authorisation status change:
26/10/2018

Reference member state:
Slovakia

Procedure number:
SK/V/0108/001

Concerned member states:
Belgium Czechia Germany Hungary Netherlands Poland

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: 3/12/2021

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