# 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
Pharmavac PHA Emulsie voor injectie
Pharmavac PHA Emulsion injectable
Pharmavac PHA Emulsion zur Injektion

**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)

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**Pharmaceutical form:**

Emulsion for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01EA

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Belgium

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**Package description:**

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

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## 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:**
23/01/2019

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

**Responsible authority:**
Federal Agency For Medicines And Health Products

**Authorisation number:**
BE-V538355

**Date of authorisation status change:**
23/01/2019

**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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.