

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:

Poland

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:

8/02/2019

Manufacturing sites for batch release:

Pharmagal Bio spol. s r.o.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2838

Date of authorisation status change:

8/02/2019

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.

Combined File of all Documents