# RP Vacc emulsion for injection for pigeons

Authorised

- Pigeon paramyxovirus 1, strain 988M, Inactivated
- Pigeon rotavirus A, strain Ro/D, Inactivated

## Product identification

**Medicine name:**
RP Vacc emulsion for injection for pigeons
RP Vacc Emulsja do wstrzykiwań

**Active substance:**
Pigeon paramyxovirus 1, strain 988M, Inactivated
Pigeon rotavirus A, strain Ro/D, Inactivated

**Target species:**
Pigeon

**Route of administration:**
Intramuscular use
Subcutaneous use

## Product details

**Active substance and strength:**
Pigeon paramyxovirus 1, strain 988M, Inactivated
6.47 log2 haemagglutination inhibiting unit(s) / 0.30 millilitre(s)

Pigeon rotavirus A, strain Ro/D, Inactivated
52.20 enzyme-linked immunosorbent assay unit / 0.30 millilitre(s)

**Pharmaceutical form:**
Emulsion for injection

**Withdrawal period by route of administration:**
**Intramuscular use:**

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

**Subcutaneous use:**

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

**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:**
Cardboard box 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:**
30/12/2021

**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:**
3152

**Date of authorisation status change:**
30/12/2021

**Reference member state:**
Slovakia

**Procedure number:**
SK/V/0110/001

**Concerned member states:**
Belgium Czechia Germany Hungary Netherlands Poland Portugal Romania

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.