Colombovac PMV/POX

Authorised

- Newcastle disease virus, strain La Sota, Inactivated
- Pigeonpox virus, strain DD, Live

Product identification

Medicine name:
Colombovac PMV/POX

Active substance:
Newcastle disease virus, strain La Sota, Inactivated

Pigeonpox virus, strain DD, Live

Target species:
Pigeon

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Newcastle disease virus, strain La Sota, Inactivated
19.90 antigen unit(s) / 0.20 millilitre(s)

Pigeonpox virus, strain DD, Live
103.50 tissue culture infective dose 50 / 0.20 millilitre(s)

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Pigeon**
  - Meat and offal. 0 day — Pombos a partir das 6 semanas de idade

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01EH01

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

Authorisation status:
Valid

Authorised in:
Portugal

Package description:
Available only in Portuguese

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:
Zoetis Portugal Lda.

Marketing authorisation date:
1/06/1999

Manufacturing sites for batch release:
Zoetis Manufacturing & Research Spain S.L.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
597/98 DGV

Date of authorisation status change:
1/11/2016

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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