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Chevipok

Authorised

- Pigeonpox virus, strain NJ, Live

Product identification

Medicine name:
Chevipok

Active substance:
Pigeonpox virus, strain NJ, Live

Target species:
Pigeon

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Pigeonpox virus, strain NJ, Live
1000000.00 50% Embryo Infective Dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for oral suspension

Withdrawal period by route of administration:

Subcutaneous use:

- **Pigeon**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01ED01

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

Authorisation status:
Valid

Authorised in:
Germany

Package description:
Available only in German
Available only in German
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Chevita Tierarzneimittel GmbH

Marketing authorisation date:
24/06/2011

Manufacturing sites for batch release:

Chevita Tierarzneimittel GmbH

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.11530.01.1

Date of authorisation status change:
1/04/2016

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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