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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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