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MD VAC LYO LYOPHILISAT POUR SUSPENSION INJECTABLE POUR POULES

Authorised

- Turkey herpesvirus, strain FC-126 (cell-associated), Live

Product identification

Medicine name:

MD VAC LYO LYOPHILISAT POUR SUSPENSION INJECTABLE POUR POULES

Active substance:

Turkey herpesvirus, strain FC-126 (cell-associated), Live

Target species:

Chicken (layer hen)

Chicken (one day-old chick)

Chicken (for reproduction)

Chicken (broiler)

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Turkey herpesvirus, strain FC-126 (cell-associated), Live
1000.00 plaque forming unit / 1.00 Dose

Pharmaceutical form:

Lyophilisate for suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

•

Chicken (layer hen)

- All relevant tissues. 0 day

•

Chicken (one day-old chick)

- All relevant tissues. 0 day

•

Chicken (for reproduction)

- All relevant tissues. 0 day

•

Chicken (broiler)

- All relevant tissues. 0 day

Subcutaneous use:

•

Chicken (layer hen)

- All relevant tissues. 0 day

•

Chicken (one day-old chick)

- All relevant tissues. 0 day

•

Chicken (for reproduction)

- All relevant tissues. 0 day

•

Chicken (broiler)

- All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Zoetis France

Marketing authorisation date:

10/08/1981

Manufacturing sites for batch release:

Zoetis Belgium

Zoetis Manufacturing & Research Spain S.L.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/6625954 1/1981

Date of authorisation status change:

10/08/2011

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 and Labelling

This document does not exist in this language (English). You can find it in another language below.