

ZYLEXIS, Lyofilizát a rozpouštědlo pro injekční suspenzi

Authorised

- Orf virus, strain D1701, Inactivated

Product identification

Medicine name:

ZYLEXIS, Lyofilizát a rozpouštědlo pro injekční suspenzi

Active substance:

Orf virus, strain D1701, Inactivated

Target species:

Dog

Cat

Horse

Cattle

Pig

Route of administration:

Subcutaneous use

Intramuscular use

Product details

Active substance and strength:

Orf virus, strain D1701, Inactivated

1.00 relative potency / 1.00 Dose

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

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

-

Dog

-

Cat**Intramuscular use:**

-

Horse

- Meat and offal. 0 day

- Milk. 0 hour

-

Cattle

- Meat and offal. 0 day

- Milk. 0 hour

-

Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QL03AX

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Available in:

Czechia

Package description:

Available only in [Czech](#)

Available only in [Czech](#)

Available only in [Czech](#)

Available only in [Czech](#)

Available only in [Czech](#)

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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 Ceska Republika s.r.o.

Marketing authorisation date:

30/10/2003

Manufacturing sites for batch release:

Zoetis Belgium

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/079/03-C

Date of authorisation status change:

30/10/2003

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

Documents

Summary of Product Characteristics

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Package Leaflet

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Labelling

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

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