

Neoleish (--) - Nasal spray, solution

Authorised

- Bacterial DNA plasmid containing LACK gene from *Leishmania infantum*

Product identification

Medicine name:

Neoleish (--) - Nasal spray, solution

Active substance:

Bacterial DNA plasmid containing LACK gene from *Leishmania infantum*

Target species:

Dog

Route of administration:

Nasal use

Product details

Active substance and strength:

Bacterial DNA plasmid containing LACK gene from *Leishmania infantum*

Presentation_strength:212.5-250 µg Index:0

Pharmaceutical form:

Nasal spray, solution

Withdrawal period by route of administration:

Nasal use:

-

Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AX

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Vial (glass), Package_size:10 vials, Content:10 x 1 ml

Packaging:Vial (glass), Package_size:1 vial, Content:1 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

CZ Vaccines S.A.U.

Marketing authorisation date:

20/12/2022

Manufacturing sites for batch release:

CZ VETERINARIA, S.A.

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

30/05/2024

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

Documents

Combined File of all Documents

English (PDF)

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