Nobivac Lepto, süstesuspensioon koertele

Not authorised

- Leptospira interrogans, serovar Copenhageni, strain 820K, Inactivated
- Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated

Product identification

Medicine name:
Nobivac Lepto, süstesuspensioon koertele

Active substance:
Leptospira interrogans, serovar Copenhageni, strain 820K, Inactivated

Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Leptospira interrogans, serovar Copenhageni, strain 820K, Inactivated
1500.00 antigen unit(s) / 1.00 millilitre(s)

Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated
1900.00 antigen unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AB01

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

Authorisation status:
Surrendered

Authorised in:
Estonia

Package description:
Available only in Estonian
Available only in Estonian

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:
Intervet International B.V.

Marketing authorisation date:
3/04/2003

Manufacturing sites for batch release:
Intervet Nederland B.V.

Responsible authority:

State Agency Of Medicines

Authorisation number:
1137

Date of authorisation status change:
29/04/2024

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.