EURICAN DAP LYOPHILISATE AND SOLVENT FOR SUSPENSION FOR INJECTION

Not authorised

- Water for injection
- Canine distemper virus, strain BA5, Live
- Canine parvovirus, strain CAG2, Live
- Canine adenovirus 2, strain DK13, Live

Product identification

Medicine name:
EURICAN DAP LYOPHILISATE AND SOLVENT FOR SUSPENSION FOR INJECTION
Eurican DAP liofilizāts un šķīdinātājs suspensijas injekcijām pagatavošanai

Active substance:
Water for injection

Canine distemper virus, strain BA5, Live

Canine parvovirus, strain CAG2, Live

Canine adenovirus 2, strain DK13, Live

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Water for injection
1.00 millilitre(s) / 1.00 Dose

Canine distemper virus, strain BA5, Live
10000.00 50% cell culture infectious dose / 1.00 Dose

Canine parvovirus, strain CAG2, Live
79432.00 50% cell culture infectious dose / 1.00 Dose

Canine adenovirus 2, strain DK13, Live
316.00 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Dog**
  - Not applicable. no withdrawal period — Withdrawal period is 0 days

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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AD02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Surrendered

Authorised in:

Latvia

Package description:

Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
31/03/2016

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/MRP/16/0007

Date of authorisation status change:
28/08/2023

Reference member state:
France

Procedure number:
FR/V/0305/001

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

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

Labelling

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