

EURICAN DAP LYOPHILISATE AND SOLVENT FOR SUSPENSION FOR INJECTION

Not
authorised

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

Product identification

Medicine name:

EURICAN DAP LYOPHILISATE AND SOLVENT FOR SUSPENSION FOR INJECTION

Active substance:

Canine adenovirus 2, strain DK13, Live
Canine distemper virus, strain BA5, Live
Canine parvovirus, strain CAG2, Live

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canine adenovirus 2, strain DK13, Live
2.50 log₁₀ 50% cell culture infectious dose / 1.00 Dose
Canine distemper virus, strain BA5, Live
4.00 log₁₀ 50% cell culture infectious dose / 1.00 Dose
Canine parvovirus, strain CAG2, Live
4.90 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AD02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Hungary

Package description:

Plastic box of 10 vials of lyophilisate (1 dose)
Plastic box of 50 vials of lyophilisate (1 dose)
Plastic box of 10 vials of solvent (1 ml)
Plastic box of 50 vials of solvent (1 ml)
Plastic box of 10 vials of lyophilisate (1 dose) and 10 vials of solvent (1 ml)
Plastic box of 50 vials of lyophilisate (1 dose) and 50 vials of solvent (1 ml)

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:

10/03/2016

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

3722/X/16 NÉBIH ÁTI

Date of authorisation status change:

10/03/2016

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

eu-puar-frv0305001-mr-rpe791-en.pdf