EURICAN DAP-L, liofilizzato e solvente per sospensione iniettabile per cani

Not authorised

- Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated
- Leptospira interrogans, serovar Canicola, Inactivated
- Canine parvovirus, Live
- Canine adenovirus 2, Live

Product identification

Medicine name:
EURICAN DAP-L, liofilizzato e solvente per sospensione iniettabile per cani

Active substance:
Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

Leptospira interrogans, serovar Canicola, Inactivated

Canine parvovirus, Live

Canine adenovirus 2, Live

Target species:
Dog

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated
1.00 European Pharmacopoeia Unit(s) / 1.00 Vial

Leptospira interrogans, serovar Canicola, Inactivated
1.00 European Pharmacopoeia Unit(s) / 1.00 Vial

Canine parvovirus, Live
4.90 log10 50% cell culture infectious dose / 1.00 Dose

Canine adenovirus 2, Live
2.50 log10 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AJ09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Italy

Package description:

Available only in Italian
Available only in Italian
Available only in Italian

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:
Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

Marketing authorisation date:
9/06/1989

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France SCS

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
9/06/2019

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

Documents

Combined File of all Documents

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