Eurican Herpes 205 (--) - Powder and solvent for emulsion for injection

Authorised

- Canine herpesvirus, strain F205, glycoproteins

Product identification

Medicine name:
Eurican Herpes 205 (--) - Powder and solvent for emulsion for injection

Active substance:
Canine herpesvirus, strain F205, glycoproteins

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Canine herpesvirus, strain F205, glycoproteins

Presentation_strength:0.3 µg to 1.75 µg Reference:Hse Comments:Antigens (amount expressed in µg of gB glycoproteins) Index:0

Pharmaceutical form:
Powder and solvent for emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AA

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)

Available in:
Austria , Bulgaria , Czechia , France , Hungary , Poland , Slovakia , Spain

Package description:
Packaging:Powder: bottle (glass); solvent: bottle (glass), Package_size:Powder: 10 bottles; solvent 10 bottles, Content:Powder: 1 dose; solvent: 1 ml
Packaging:Powder: bottle (glass); solvent: bottle (glass), Package_size:Powder: 50 bottles; solvent: 50 bottles, Content:Powder: 1 ml; solvent: 1 ml
Packaging:Powder: bottle (glass); solvent: bottle (glass), Package_size:Powder: 1 bottle; solvent: 1 bottle, Content:Powder: 1 dose; solvent: 1 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
26/03/2001

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
European Commission

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

Date of authorisation status change:
26/03/2001

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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