FIXR MYC-VAC

Authorised

- Mycoplasma gallisepticum, strains MG-NEV40 and MG-NEV45, Inactivated

Product identification

Medicine name:
FIXR MYC-VAC

Active substance:
Mycoplasma gallisepticum, strains MG-NEV40 and MG-NEV45, Inactivated

Target species:
Chicken

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Mycoplasma gallisepticum, strains MG-NEV40 and MG-NEV45, Inactivated
40.00 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AB03

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

Authorisation status:
Valid

Authorised in:
Portugal

Package description:
Polystyrene box: 10 x 250 ml polypropylene bottles (Ph. Eur.) which are closed with elastomer stoppers (Ph.Eur.) and sealed with aluminium caps.
Carton box: 1 x 250 ml polypropylene bottle (Ph. Eur.) which are closed with elastomer stoppers (Ph.Eur.) and sealed with aluminium caps.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Similar biological application (Article 13(4) of Directive No 2001/82/EC)

Marketing authorisation holder:
Kernfarm B.V.

Marketing authorisation date:
22/02/2024

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
1627/01/24 DIVPT

Date of authorisation status change:
22/02/2024

Reference member state:
Netherlands

Procedure number:
NL/V/0363/001

Concerned member states:
Belgium Czechia Germany Hungary Portugal Romania Spain

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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