

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:

Germany

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:

6/03/2026

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Paul-Ehrlich-Institut

Authorisation number:

PEI.V.12280.01.1

Date of authorisation status change:

6/03/2026

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

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