Ingelvac Ery emulsion for injection for pigs

Authorised

- Erysipelothrix rhusiopathiae, serotype 2, strain SE-9, Inactivated

Product identification

Medicine name:

Ingelvac Ery emulsion for injection for pigs
INGELVAC ERY emulsione iniettabile per suini

Active substance:

Erysipelothrix rhusiopathiae, serotype 2, strain SE-9, Inactivated

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Erysipelothrix rhusiopathiae, serotype 2, strain SE-9, Inactivated
61.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AB03

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

Authorisation status:
Valid

Authorised in:
Italy

Package description:
cardboard box containing 1 vial of 100 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:
Laboratorios Syva S.A.

Marketing authorisation date:
7/02/2022

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
Ministry Of Health

Authorisation number:
105518

Date of authorisation status change:

7/02/2022

Reference member state:

Spain

Procedure number:

ES/V/0395/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

Package Leaflet

Labelling