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ERYSIN emulsion for injection

Authorised

- Erysipelothrix rhusiopathiae, Inactivated

Product identification

Medicine name:
ERYSIN emulsion for injection

Active substance:
Erysipelothrix rhusiopathiae, Inactivated

Target species:
Pig

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Erysipelothrix rhusiopathiae, Inactivated
100.00 Colony forming 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:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

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:
Bioveta a.s.

Marketing authorisation date:
13/05/1997

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1720

Date of authorisation status change:

13/05/1997

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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