# Eryseng (--) - Suspension for injection

Authorised

- Erysipelothrix rhusiopathiae, strain R32E11, Inactivated

## Product identification

**Medicine name:**
Eryseng (--) - Suspension for injection

**Active substance:**
Erysipelothrix rhusiopathiae, strain R32E11, Inactivated

**Target species:**
Pig

**Route of administration:**
Intramuscular use

## Product details

**Active substance and strength:**
Erysipelothrix rhusiopathiae, strain R32E11, Inactivated

Presentation_strength:ELISA > 3.34 log2 IE50%/dose Reference:Ph Eur Index:0

**Pharmaceutical form:**
Suspension 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:**
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)

**Package description:**
Packaging:Bottle (PET), Package_size:1 bottle, Content:250 ml (125 doses)
Packaging:Bottle (PET), Package_size:1 bottle, Content:100 ml (50 doses)
Packaging:Bottle (PET), Package_size:1 bottle, Content:50 ml (25 doses)
Packaging:Bottle (PET), Package_size:1 bottle, Content:20 ml (10 doses)
Packaging:Vial (glass), Package_size:1 vial, Content:100 ml (50 doses)
Packaging:Vial (glass), Package_size:1 vial, Content:50 ml (25 doses)
Packaging:Vial (glass), Package_size:1 vial, Content:20 ml (10 doses)

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Laboratorios Hipra, S.A.

**Marketing authorisation date:**
4/07/2014

**Manufacturing sites for batch release:**

Laboratorios Hipra S.A.

**Responsible authority:**
European Commission

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

**Date of authorisation status change:**
4/07/2014

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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