Suvaxyn® Parvo/E

Not authorised

- Erysipelothrix rhusiopathiae, serotype 2, Inactivated
- Porcine parvovirus, strain S-80, Inactivated

Product identification

Medicine name:
Suvaxyn® Parvo/E

Active substance:
Erysipelothrix rhusiopathiae, serotype 2, Inactivated

Porcine parvovirus, strain S-80, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Erysipelothrix rhusiopathiae, serotype 2, Inactivated
13.50 relative potency / 2.00 millilitre(s)

Porcine parvovirus, strain S-80, Inactivated
94.10 haemagglutination inhibiting unit(s) / 2.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AL01

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

Authorisation status:
Revoked

Authorised in:
Portugal

Package description:
Available only in Portuguese
Available only in Portuguese

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 8(3) of Directive No 2001/83/EC)

Marketing authorisation holder:
Zoetis Portugal Lda.

Marketing authorisation date:
1/01/2022

Manufacturing sites for batch release:
Zoetis Portugal Lda.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
UPD 1.6.1-4 JSON NAP Chpt2 C2 Mandatory 889/01/15RIVPT

Date of authorisation status change:
1/04/2022

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