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Suvaxyn Circo+MH RTU (--) Emulsion for injection

Authorised

- Mycoplasma hyopneumoniae, strain P-5722-3, Inactivated
- Porcine circovirus type 2, ORF2 capsid protein

Product identification

Medicine name:
Suvaxyn Circo+MH RTU (--) Emulsion for injection

Active substance:
Mycoplasma hyopneumoniae, strain P-5722-3, Inactivated
Porcine circovirus type 2, ORF2 capsid protein

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Mycoplasma hyopneumoniae, strain P-5722-3, Inactivated
Presentation_strength:1.5 - 3.8 RP Reference:Hse Index:0
Porcine circovirus type 2, ORF2 capsid protein

Presentation_strength:2.3-12.4 RP Reference:Hse Index:1

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AL08

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)

Available in:

Belgium , Cyprus , Denmark , Germany , Greece , Italy , Luxembourg , Netherlands , Poland , Spain , Sweden

Package description:

Packaging:vial (HDPE), Package_size:1 vial, Content:250 ml

Packaging:vial (HDPE), Package_size:1 vial, Content:50 ml

Packaging:vial (HDPE), Package_size:1 vial, Content:100 ml

Packaging:vial (HDPE), Package_size:10 vials, Content:100 ml

Packaging:vial (HDPE), Package_size:4 vials, Content:250 ml

Packaging:vial (HDPE), Package_size:10 vials, Content:50 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:
Zoetis Belgium

Marketing authorisation date:
6/11/2015

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
European Commission

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

Date of authorisation status change:
6/11/2015

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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