ReproCyc ParvoFLEX (--) - Suspension for injection

Authorised

- Porcine parvovirus, strain 27a, VP2 protein

Product identification

Medicine name:
ReproCyc ParvoFLEX (--) - Suspension for injection

Active substance:
Porcine parvovirus, strain 27a, VP2 protein

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Porcine parvovirus, strain 27a, VP2 protein

Presentation_strength:≥1.0 RP Reference:Hse Index:0

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AA02

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 , France , Luxembourg , Spain

Package description:

Packaging:Bottle (HDPE), Package_size:12 bottles, Content:20 ml (10 doses)
Packaging:Bottle (HDPE), Package_size:1 bottle, Content:20 ml (10 doses)
Packaging:Bottle (HDPE), Package_size:1 bottle, Content:100 ml (50 doses)
Packaging:Bottle (HDPE), Package_size:12 bottles, Content:200 ml (100 doses)
Packaging:Bottle (HDPE), Package_size:12 bottles, Content:100 ml (50 doses)
Packaging:Bottle (HDPE), Package_size:1 bottle, Content:200 ml (100 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:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
26/04/2019

Manufacturing sites for batch release:
Boehringer Ingelheim Vetmedica GmbH

Responsible authority:
European Commission

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

Date of authorisation status change:
26/04/2019

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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