Porcilis PCV M Hyo ID (--) + (--) - Emulsion for injection

Authorised

- Porcine circovirus type 2, ORF2 capsid protein
- Mycoplasma hyopneumoniae, strain J, Inactivated

Product identification

Medicine name:
Porcilis PCV M Hyo ID (--) + (--) - Emulsion for injection

Active substance:
Porcine circovirus type 2, ORF2 capsid protein

Mycoplasma hyopneumoniae, strain J, Inactivated

Target species:
Pig

Route of administration:
Intradermal use

Product details

Active substance and strength:
Porcine circovirus type 2, ORF2 capsid protein

Presentation_strength:751.4 AU Reference:HSE Index:0
Mycoplasma hyopneumoniae, strain J, Inactivated

Presentation_strength:0.72 AU Reference:HSE Index:1

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intradermal 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:

Portugal

Package description:

Packaging:Vial (PET), Package_size:1 vial, Content:20 ml
Packaging:Vial (PET), Package_size:1 vial, Content:10 ml
Packaging:Vial (glass), Package_size:10 vials, Content:100 ml
Packaging:Vial (PET), Package_size:1 vial, Content:40 ml
Packaging:Vial (glass), Package_size:1 vial, Content:10 ml
Packaging:Vial (PET), Package_size:10 vials, Content:400 ml
Packaging:Vial (PET), Package_size:10 vials, Content:100 ml
Packaging:Vial (PET), Package_size:10 vials, Content:200 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:
INTERVET INTERNATIONAL B.V.

Marketing authorisation date:
30/08/2024

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
European Commission

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

Date of authorisation status change:
30/08/2024

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

Documents

Combined File of all Documents

English (PDF)

Published on: 9/10/2024

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