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Mhyosphere PCV ID (--) - Emulsion for injection

Authorised

- Mycoplasma hyopneumoniae, strain Nexhyon, expressing capsid protein of porcine circovirus type 2A, Inactivated

Product identification

Medicine name:
Mhyosphere PCV ID (--) - Emulsion for injection

Active substance:
Mycoplasma hyopneumoniae, strain Nexhyon, expressing capsid protein of porcine circovirus type 2A, Inactivated

Target species:
Pig

Route of administration:
Intradermal use

Product details

Active substance and strength:
Mycoplasma hyopneumoniae, strain Nexhyon, expressing capsid protein of porcine circovirus type 2A, Inactivated

Presentation_strength:RP* ≥1.3 Reference:HSE Index:0

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)

Package description:

Packaging:Vial (PET), Package_size:1 vial, Content:20 ml (100 doses)
Packaging:Vial (PET), Package_size:1 vial, Content:10 ml (50 doses)
Packaging:Vial (PET), Package_size:1 vial, Content:25 ml (125 doses)
Packaging:Vial (PET), Package_size:1 vial, Content:50 ml (250 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:
18/09/2020

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:
18/09/2020

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: 7/10/2026
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