

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:50 ml (250 doses)

Packaging:Vial (PET), Package_size:1 vial, Content:25 ml (125 doses)

Packaging:Vial (PET), Package_size:1 vial, Content:10 ml (50 doses)

Packaging:Vial (PET), Package_size:1 vial, Content:20 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:

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)

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