PORCILIS M HYO, SUSPENSION FOR INJECTION

Not authorised

- Mycoplasma hyopneumoniae, strain 11, Inactivated

Product identification

Medicine name:
PORCILIS M HYO, SUSPENSION FOR INJECTION

Active substance:
Mycoplasma hyopneumoniae, strain 11, Inactivated

Target species:
Pig (for fattening)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Mycoplasma hyopneumoniae, strain 11, Inactivated
7.00 log2 antibody unit(s) / 1.00 unit(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig (for fattening)**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Surrendered

Authorised in:

Austria

Package description:

Cardboard boxes with 1 PET vial of 20 ml (10 doses)
Cardboard boxes with 5 PET vials of 20 ml (10 doses)
Cardboard boxes with 10 PET vials of 20 ml (10 doses)
Cardboard boxes with 1 PET vial of 50 ml (25 doses)
Cardboard boxes with 5 PET vials of 50 ml (25 doses)
Cardboard boxes with 10 PET vials of 50 ml (25 doses)
Cardboard boxes with 1 PET vial of 100 ml (50 doses)
Cardboard boxes with 5 PET vials of 100 ml (50 doses)
Cardboard boxes with 10 PET vials of 100 ml (50 doses)
Cardboard boxes with 1 PET vial of 200 ml (100 doses)
Cardboard boxes with 5 PET vials of 200 ml (100 doses)
Cardboard boxes with 10 PET vials of 200 ml (100 doses)
Cardboard boxes with 1 PET vial of 250 ml (125 doses)
Cardboard boxes with 5 PET vials of 250 ml (125 doses)
Cardboard boxes with 10 PET vials of 250 ml (125 doses)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet Ges.m.b.H.

Marketing authorisation date:
23/03/2006

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

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-20284

Date of authorisation status change:
12/11/2024

Reference member state:
France

Procedure number:
FR/V/0158/001

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

Documents

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.