INGELVAC MYCOFLEX SUSPENSION FOR INJECTION FOR PIGS

Authorised

- Mycoplasma hyopneumoniae, strain J, Inactivated

Product identification

Medicine name:
INGELVAC MYCOFLEX SUSPENSION FOR INJECTION FOR PIGS
Ingelvac MycoFLEX suspensie voor injectie voor varkens

Active substance:
Mycoplasma hyopneumoniae, strain J, Inactivated

Target species:
Pig (for fattening)
Pig (for reproduction)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Mycoplasma hyopneumoniae, strain J, Inactivated
1.00 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AB13

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Cardboard box with 1 high density polyethylene bottle of 10 ml (10 doses)
Cardboard box with 12 high density polyethylene TwistPak bottles of 250 ml (250 doses)
Cardboard box with12 high density polyethylene TwistPak bottles of 100 ml (100 doses)
Cardboard box with 12 high density polyethylene TwistPak bottles 50 ml (50 doses)
Cardboard box with 12 high density polyethylene TwistPak bottles of 10 ml (10 doses)
Cardboard box with 1 density polyethylene TwistPak bottle of 250 ml (250 doses in 250 ml) vaccine
Cardboard box with 1 density polyethylene TwistPak bottle of 100 ml (100 doses)
Cardboard box with 1 density polyethylene TwistPak bottle of 50 ml (50 doses)
Cardboard box with 1 density polyethylene TwistPak bottle of 10 ml (10 doses)

Cardboard box with 12 high density polyethylene bottles of 250 ml (250 doses)
Cardboard box with 12 high density polyethylene bottles of 100 ml (100 doses)
Cardboard box with 12 high density polyethylene bottles of 50 ml (50 doses)
Cardboard box with 12 high density polyethylene bottles of 10 ml (10 doses)
Cardboard box with 1 high density polyethylene bottle of 250 ml (250 doses)
Cardboard box with 1 high density polyethylene bottle of 100 ml (100 doses)
Cardboard box with 1 high density polyethylene bottle of 50 ml (50 doses)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
27/04/2009

Manufacturing sites for batch release:
Boehringer Ingelheim Vetmedica GmbH

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 104086

Date of authorisation status change:
20/06/2022

Reference member state:
France

Procedure number:
FR/V/0203/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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