Porcilis M Hyo ID ONCE emulsion for injection for pigs

Authorised

- Mycoplasma hyopneumoniae, strain 11, Inactivated

Product identification

Medicine name:
Porcilis M Hyo ID ONCE emulsion for injection for pigs
Porcilis M Hyo ID ONCE injektionsvæske, emulsion

Active substance:
Mycoplasma hyopneumoniae, strain 11, Inactivated

Target species:
Pig

Route of administration:
Intradermal use

Product details

Active substance and strength:
Mycoplasma hyopneumoniae, strain 11, Inactivated

Pharmaceutical form:
Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AB13

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

Authorisation status:
Valid

Authorised in:
Denmark

Available in:
Denmark

Package description:
Cardboard box with 10 PET vials of 20 ml (100 doses)
Cardboard box with 5 PET vials of 20 ml (100 doses)
Cardboard box with 1 PET vial of 20 ml (100 doses)
Cardboard box with 10 glass vials of 20 ml (100 doses)
Cardboard box with 10 glass vials of 10 ml (50 doses)
Cardboard box with 5 glass vials of 20 ml (100 doses)
Cardboard box with 5 glass vials of 10 ml (50 doses)
Cardboard box with 1 glass vial of 20 ml (100 doses)
Cardboard box with 1 glass vial of 10 ml (50 doses)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:

2/11/2011

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

Responsible authority:
Danish Medicines Agency

Authorisation number:
47083

Date of authorisation status change:
2/11/2011

Reference member state:
Hungary

Procedure number:
HU/V/0109/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia France Germany Greece Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway 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

Summary of Product Characteristics

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

Combined File of all Documents