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M+PAC

Authorised

- Mycoplasma hyopneumoniae, Inactivated

Product identification

Medicine name:
M+PAC

Active substance:
Mycoplasma hyopneumoniae, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Mycoplasma hyopneumoniae, Inactivated
1.47 relative unit(s) / 1.00 millilitre(s)

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:
Ireland

Package description:
Box of 10 bottle of 200 ml
Box of 5 bottle of 200 ml
Box of 2 bottle of 200 ml
Box of 1 bottle of 200 ml
Box of 1 bottle of 50 ml
Box of 10 bottle of 100 ml
Box of 5 bottle of 100 ml
Box of 2 bottle of 100 ml
Box of 1 bottle of 100 ml
Box of 10 bottles of 50 ml
Box of 5 bottles of 50 ml
Box of 2 bottles of 50 ml

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 (Ireland) Limited

Marketing authorisation date:
12/09/2002

Manufacturing sites for batch release:
Burgwedel Biotech GmbH

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10996/236/001

Date of authorisation status change:
12/09/2002

Reference member state:
Hungary

Procedure number:
HU/V/0140/001

Concerned member states:
Austria Belgium Cyprus Czechia Denmark Estonia Germany Greece Ireland Italy Latvia Lithuania Luxembourg Poland Portugal 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