M+PAC

Authorised

- Mycoplasma hyopneumoniae, Inactivated

Product identification

Medicine name:
M+PAC
ThoroVAX vet., injekcinė emulsija

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

Withdrawal period by route of administration:
Intramuscular use:

- **Pig**
  - Meat and offal. 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:
Lithuania

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 International B.V.

Marketing authorisation date:
16/10/2005

Manufacturing sites for batch release:
Burgwedel Biotech GmbH

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/05/1690/001-003

Date of authorisation status change:
22/05/2007

Reference member state:
Hungary

Procedure number:
HU/V/0140/001/MR

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

RV1690.pdf
