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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:
Lithuania

Available 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:
1/04/2024

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)

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Documents

Combined File of all Documents

English (PDF)
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