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M+PAC

Not 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:
Surrendered

Authorised in:
Denmark

Package description:
Box of 1 bottle of 200 ml
Box of 5 bottle of 200 ml
Box of 10 bottle of 200 ml
Box of 1 bottle of 50 ml
Box of 5 bottle of 100 ml
Box of 2 bottle of 200 ml
Box of 5 bottles of 50 ml
Box of 10 bottles of 50 ml
Box of 1 bottle of 100 ml
Box of 2 bottle of 100 ml
Box of 10 bottle of 100 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:
3/07/2002

Manufacturing sites for batch release:
Burgwedel Biotech GmbH

Responsible authority:
Danish Medicines Agency

Authorisation number:
33449

Date of authorisation status change:
27/10/2025

Reference member state:
Hungary

Procedure number:
HU/V/0140/001

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

Documents

Summary of Product Characteristics

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Combined File of all Documents