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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:
Revoked

Authorised in:
Portugal

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:
MSD Animal Health Lda.

Marketing authorisation date:
1/10/2025

Manufacturing sites for batch release:
Burgwedel Biotech GmbH

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
R706/02 DGV

Date of authorisation status change:
1/11/2014

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

Combined File of all Documents

English (PDF)
Published on: 22/08/2025
Updated on: 13/03/2026
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