

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

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents