

# M+PAC

Authorised

- Mycoplasma hyopneumoniae, Inactivated

## Product identification

**Medicine name:**

M+PAC

M + PAC Emulsie voor injectie

M + PAC Emulsion injectable

M + PAC Emulsion zur Injektion

**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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI09AB13

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Belgium

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

28/11/2005

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**Manufacturing sites for batch release:**

Burgwedel Biotech GmbH

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**Responsible authority:**

FAMHP

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

28/11/2005

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**Reference member state:**

Hungary

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**Procedure number:**

HU/V/0140/001

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**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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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Package Leaflet

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

### Summary of Product Characteristics

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

### Labelling

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

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