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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:

Luxembourg

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:

28/11/2005

Manufacturing sites for batch release: Burgwedel Biotech GmbH
Responsible authority: Ministry Of Health And Social Security
Authorisation number: V 817/02/08/0734
Date of authorisation status change: 20/10/2025
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
Combined File of all Documents