M+PAC

Authorised

- Mycoplasma hyopneumoniae, Inactivated

Product identification

Medicine name:
M+PAC
M+Pac - Emulsion zur Injektion für Schweine

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**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AB13

Legal status of supply:
Medicinal product on medical prescription for non-renewable delivery

Authorisation status:
Valid

Authorised in:
Austria

Package description:
Available only in German
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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 Ges.m.b.H.

Marketing authorisation date:
10/07/2002

Manufacturing sites for batch release:
Burgwedel Biotech GmbH

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-20248

Date of authorisation status change:
14/11/2006

Reference member state:
Hungary

Procedure number:
HU/V/0140/001

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)

To consult adverse reactions on veterinary medicinal products please go to 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.
